

PROJECT NOTE

To: Jard Company Inc. Hazard Ranking System Project File

From: John Burton, Weston Solutions, Inc. (WESTON®), Superfund Technical Assessment

and Response Team III (START)

Thru: Mr. John F. Kelly, Project Leader, START

Date: 2 October 2013

RE: Source, Surface Soil, and Sediment Sample Sample-Adjusted Contract Required

Quantitation Limit Calculations and Form Is

Case 43392; SDG A4B24

TDD No. 13-09-0001; Task No. 0904-48; DC No. A-6871

Introduction

The following Project Note describes the sample-adjusted Contract Required Quantitation Limit (CRQL) calculations for polychlorinated biphenyls (PCBs) analytical results of 19 source samples collected from the Jard Company Inc. property located in Bennington, Bennington County, Vermont. The samples were collected by Weston Solutions, Inc. (WESTON®), Superfund Technical Assessment and Response Team III (START) for the purpose of performing a Site Reassessment in support of a U.S. Environmental Protection Agency (EPA) Hazard Ranking System (HRS)/National Priorities List (NPL) Documentation Record. The analytical data were validated at a Tier II level according to Region I EPA-NE Data Validation Functional Guidelines for Evaluating Environmental Analyses and the USEPA CLP National Functional Guidelines for Superfund Organic Methods.

Table 1 of this Project Note summarizes the validated analytical results for PCB analysis. Table 2 of this Project Note reports the sample-adjusted CRQL for each sample, which is either the CRQL or a raised value due to the dilution factor, percent solids, sample volume, and/or final volume. Tables 1 and 2 are included in *Attachment A* of this Project Note. The memorandum detailing the original validated results is included in *Attachment B* of this Project Note.

Copies of the pertinent Form I's have been included in *Attachment C* of this Project Note. CRQLs are listed in the *USEPA Contract Laboratory Program Statement of Work for Organics Analysis, Multi-media Multi-concentration, SOM01.1*, and the *Modifications Updating SOM01.1* to *SOM01.2*, for PCBs, the pertinent portions of which are included in *Attachment D* of this Project Note.



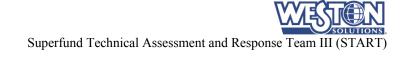
Sample-adjusted CRQL Determination for Soil/Source and Sediment Samples:

The sample-adjusted CRQLs were calculated as follows: the PCB sample-adjusted CRQLs [in micrograms per kilogram ($\mu g/Kg$)] was calculated by multiplying the CRQL (in $\mu g/Kg$) for the substance by the method extraction weight [30 grams (g) nominally], dividing this result by the dry weight extracted (in g), and multiplying this result by the dilution factor. The dry weight extracted (in g) is calculated by multiplying the percent solids, expressed as a decimal, of the sample (100% - percent moisture) by the wet weight extracted (in g) of the sample. The percent moisture, wet weight extracted, and dilution factors are reported on the Form I for the sample.

Sample-adjusted CRQL (
$$\mu g/Kg$$
) =
$$\frac{[CRQL \times 30g]}{[\%S \times W_W]} / \times DF$$

 $CRQL = in \mu g/Kg$ %S = Percent Solids (in decimal form)

 W_W = wet weight extracted (g) DF = Dilution Factor



Attachment A

Tables

HRS Reference #81 Page 3 of 126

DATA SUMMARY TABLE 1 AROCLOR IN SOIL ANALYSIS $\mu g/Kg$

CONSULTING GROUP

	SAMI	PLE NUMBER	A4B24	A4B25	A4B26	A4B27	A4B28	A4B29	A4B33
	SAMPLE LOCATION		SO-07	SO-14	SO-200	SO-21	SO-22	SO-23	SO-62
	STATIO	N LOCATION	JCS-008	JCS-015	JCS-475	JCS-024	JCS-025	JCS-026 E1902-06	JCS-076 E1902-09
	LABORATO	RY NUMBER	E1902-01	E1902-02	E1902-03	E1902-04	E1902-05		
COMPOUND	MDL	CRQL							
Aroclor-1016	2.6	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1221	7.8	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1232	1.3	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1242	6.2	33	41 U	120	130	110 J	140	36 U	180
Aroclor-1248	2.7	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1254	3.2	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1260	3.2	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1262	14	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1268	6.6	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
	DILUT	ION FACTOR	1.0	1.0	1.0	1.0	1.0	1.0	1.0
	DATE SAMPLED		4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/4/2013
DATE EXTRACTED		EXTRACTED	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013
DATE ANALYZED		4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013	
	SAMPLE WEI	GHT (GRAMS)	30	30.1	30.0	30.0	30.0	30.0	30.1
		% SOLID	80.4	88.8	88.4	90.8	91.0	90.6	85.6

NOTES: μg/Kg = micrograms per Kilogram

All results are reported on a Dry Weight Basis.

MDL = Method Detection Limit

CRQL = Contract Required Quantitation Limit

U = Value is Non-Detected.

UJ = Value is Non-Detected, and Detection Limit is Estimated.

J = Value is Estimated.

R = Value is Rejected.

DATA SUMMARY TABLE 1 AROCLOR IN SOIL ANALYSIS $\mu g/Kg$

CONSULTING GROUP

	SAMPLE NUMBER		A4B35	A4B17	A4B18	A4B19	A4B20	A4B21	A4B22
	SAMPLE LOCATION		SO-65	SB-01	SB-03	SB-05	SB-06	SB-08	SB-09
	STATIO	N LOCATION	JCS-086	JCS-130	JCS-136	JCS-138	JCS-148	JCS-153	JCS-143
	LABORATO	RY NUMBER	E1902-10	902-10 E1902-13 E1902-14 E1902-15 E1902-	E1902-16	E1902-17	E1902-18		
COMPOUND	MDL	CRQL							
Aroclor-1016	2.6	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1221	7.8	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1232	1.3	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1242	6.2	33	38 U	280000 *	4800000 *	820 *	1900 *	730000 *	40000 *
Aroclor-1248	2.7	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1254	3.2	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1260	3.2	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1262	14	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1268	6.6	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
	DILUT	ION FACTOR	1.0	100 / 1000*	100 / 2000*	1 / 2*	1 / 10*	100 / 2000*	1 / 100*
	DATE SAMPLED		4/5/2013	4/1/2013	4/1/2013	4/8/2013	4/8/2013	4/8/2013	4/8/2013
DATE EXTRACTED		EXTRACTED	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013
	DATE ANALYZED		4/25/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013
	SAMPLE WEI	GHT (GRAMS)	30	30	30	30.1	30.1	30.1	30.1
		% SOLID	85.7	84.1	93.5	90.2	90.7	91.0	92.4

NOTES: μg/Kg = micrograms per Kilogram

All results are reported on a Dry Weight Basis.

MDL = Method Detection Limit

CRQL = Contract Required Quantitation Limit

U = Value is Non-Detected.

UJ = Value is Non-Detected, and Detection Limit is Estimated.

J = Value is Estimated.

R = Value is Rejected.

DATA SUMMARY TABLE 1 AROCLOR IN SOIL ANALYSIS $\mu g/Kg$

CONSULTING GROUP

	SAME	PLE NUMBER	A4B23	A4B30	A4B31	A4B32	A4B34	
	SAMPL	E LOCATION	SO-06	SO-34	SO-36	SO-46	SO-64	
	STATIO	N LOCATION	JCS-006	JCS-046	JCS-048	JCS-061	JCS-183	
	LABORATO	RY NUMBER	E1902-19	E1902-20	E1902-21	E1902-22	E1902-23	
COMPOUND	MDL	CRQL						
Aroclor-1016	2.6	33	39 U	37 U	39 U	38 U	38 U	
Aroclor-1221	7.8	33	39 U	37 U	39 U	38 U	38 U	
Aroclor-1232	1.3	33	39 U	37 U	39 U	38 U	38 U	
Aroclor-1242	6.2	33	150 J	320	1600 *	1200 *	280 J	
Aroclor-1248	2.7	33	39 U	37 U	39 U	38 U	38 U	
Aroclor-1254	3.2	33	39 U	37 U	39 U	38 U	38 U	
Aroclor-1260	3.2	33	39 U	37 U	39 U	38 U	38 U	
Aroclor-1262	14	33	39 U	37 U	39 U	38 U	38 U	
Aroclor-1268	6.6	33	39 U	37 U	39 U	38 U	38 U	
	DILUT	ION FACTOR	1.0	1.0	1 / 10*	1 / 10*	1.0	
	DA	TE SAMPLED	4/3/2013	4/4/2013	4/4/2013	4/4/2013	4/4/2013	
	DATE EXTRACTED		4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/22/2013	
	DATE ANALYZED		4/25/2013	4/25/2013	4/26/2013	4/26/2013	4/25/2013	
	SAMPLE WEI	GHT (GRAMS)	30	30	30	30	30.1	
		% SOLID	85.3	88.6	84.9	88.8	85.6	

NOTES: μg/Kg = micrograms per Kilogram

All results are reported on a Dry Weight Basis.

MDL = Method Detection Limit

CRQL = Contract Required Quantitation Limit

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UJ = Value is Non-Detected, and Detection Limit is Estimated.

J = Value is Estimated.

R = Value is Rejected.

DATA SUMMARY TABLE 2 SAMPLE ADJUSTED CRQL µg/Kg

CONSULTING GROUP

	SAME	PLE NUMBER	A4B24	A4B25	A4B26	A4B27	A4B28	A4B29	A4B33	
	SAMPLE LOCATION		SAMPLE LOC	SO-07	SO-14	SO-200	SO-21	SO-22	SO-23	SO-62
	STATIO	N LOCATION	JCS-008	JCS-015	JCS-475	JCS-024	JCS-025 JCS-026	JCS-076		
	LABORATO	RY NUMBER	E1902-01	E1902-02	E1902-03	E1902-04	E1902-05	E1902-06	E1902-09	
COMPOUND	MDL	CRQL								
Aroclor-1016	2.6	33	41	37	37	36	36	36	38	
Aroclor-1221	7.8	33	41	37	37	36	36	36	38	
Aroclor-1232	1.3	33	41	37	37	36	36	36	38	
Aroclor-1242	6.2	33	41	37	37	36	36	36	38	
Aroclor-1248	2.7	33	41	37	37	36	36	36	38	
Aroclor-1254	3.2	33	41	37	37	36	36	36	38	
Aroclor-1260	3.2	33	41	37	37	36	36	36	38	
Aroclor-1262	14	33	41	37	37	36	36	36	38	
Aroclor-1268	6.6	33	41	37	37	36	36	36	38	
	DILUT	ION FACTOR	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
DATE SAMPLED		4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/4/2013		
DATE EXTRACTED		4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013		
DATE ANALYZED		4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013		
	SAMPLE WEI	GHT (GRAMS)	30	30.1	30.0	30.0	30.0	30.0	30.1	
		% SOLID	80.4	88.8	88.4	90.8	91.0	90.6	85.6	

NOTES: μg/Kg = micrograms per Kilogram

All results are reported on a Dry Weight Basis.

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CRQL = Contract Required Quantitation Limit

U = Value is Non-Detected.

UJ = Value is Non-Detected, and Detection Limit is Estimated.

J = Value is Estimated.

R = Value is Rejected.

^{* =} Reported value is from diluted analysis.

DATA SUMMARY TABLE 2 SAMPLE ADJUSTED CRQL µg/Kg

CONSULTING GROUP

	SAMI	PLE NUMBER	A4B35	A4B17	A4B18	A4B19	A4B20	A4B21	A4B22
	SAMPLE	E LOCATION	SO-65	SB-01	SB-03	SB-05	SB-06	SB-08	SB-09
	STATIO	N LOCATION	JCS-086	JCS-086 JCS-130 JCS-136 JCS-138	JCS-148	JCS-148 JCS-153	JCS-143		
	LABORATO	RY NUMBER	E1902-10	E1902-13	E1902-14	E1902-15	E1902-16	E1902-17	E1902-18
COMPOUND	MDL	CRQL							
Aroclor-1016	2.6	33	38	3900	3500	37	36	3600	36
Aroclor-1221	7.8	33	38	3900	3500	37	36	3600	36
Aroclor-1232	1.3	33	38	3900	3500	37	36	3600	36
Aroclor-1242	6.2	33	38	39000 *	71000 *	73 *	360 *	72000 *	3600 *
Aroclor-1248	2.7	33	38	3900	3500	37	36	3600	36
Aroclor-1254	3.2	33	38	3900	3500	37	36	3600	36
Aroclor-1260	3.2	33	38	3900	3500	37	36	3600	36
Aroclor-1262	14	33	38	3900	3500	37	36	3600	36
Aroclor-1268	6.6	33	38	3900	3500	37	36	3600	36
		<u>_</u>							
	DILUT	ION FACTOR	1.0	100 / 1000*	100 / 2000*	1 / 2*	1 / 10*	100 / 2000*	1 / 100*
DATE SAMPLED		4/5/2013	4/1/2013	4/1/2013	4/8/2013	4/8/2013	4/8/2013	4/8/2013	
DATE EXTRACTED		4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	
DATE ANALYZED		4/25/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013	
	SAMPLE WEI	GHT (GRAMS)	30	30	30	30.1	30.1	30.1	30.1
		% SOLID	85.7	84.1	93.5	90.2	90.7	91.0	92.4

NOTES: μg/Kg = micrograms per Kilogram

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CONSULTING GROUP

DATA SUMMARY TABLE 2 SAMPLE ADJUSTED CRQL µg/Kg

	SAME	PLE NUMBER	A4B23	A4B30	A4B31	A4B32	A4B34	
	SAMPLE LOCATION		SO-06	SO-34	SO-36	SO-46	SO-64	
	STATIO	N LOCATION	JCS-006	JCS-046	JCS-048	JCS-061	JCS-183	
	LABORATO	RY NUMBER	E1902-19	E1902-20	E1902-21	E1902-22	E1902-23	
COMPOUND	MDL	CRQL						
Aroclor-1016	2.6	33	39	37	39	38	38	
Aroclor-1221	7.8	33	39	37	39	38	38	
Aroclor-1232	1.3	33	39	37	39	38	38	
Aroclor-1242	6.2	33	39	37	390 *	370 *	38	
Aroclor-1248	2.7	33	39	37	39	38	38	
Aroclor-1254	3.2	33	39	37	39	38	38	
Aroclor-1260	3.2	33	39	37	39	38	38	
Aroclor-1262	14	33	39	37	39	38	38	
Aroclor-1268	6.6	33	39	37	39	38	38	
	DILUT	ION FACTOR	1.0	1.0	1 / 10*	1 / 10*	1.0	
	DA	TE SAMPLED	4/3/2013	4/4/2013	4/4/2013	4/4/2013	4/4/2013	
	DATE EXTRACTED		4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/22/2013	
	DAT	E ANALYZED	4/25/2013	4/25/2013	4/26/2013	4/26/2013	4/25/2013	
	SAMPLE WEI	GHT (GRAMS)	30	30	30	30	30.1	
		% SOLID	85.3	88.6	84.9	88.8	85.6	

NOTES: μg/Kg = micrograms per Kilogram

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R = Value is Rejected.



Attachment B

Data Validation Memorandum Case No. 43392; SDG No. A4B24

HRS Reference #81 Page 10 of 126



Weston Solutions, Inc.
East Division
3 Riverside Drive
Andover, Massachusetts 01810
978-552-2100 - Fax 978-658-0700

SUPERFUND TECHNICAL ASSESSMENT AND RESPONSE TEAM EPA CONTRACT EP-W-05-042

21 August 2013 20114-081-998-0850-49 DC No. A-6844

Ms. Martha Bosworth U.S. EPA Region I - New England Emergency Planning & Response Branch 5 Post Office Square, Suite 100 Mail Code OSRR07-2 Boston, Massachusetts 02109-3912

Subject:

Case No. 43392; SDG No. A4B24

ChemTech Consulting Group (Chem)

Jard Company Inc Bennington, Vermont

AROCLOR: 19/Soil/A4B24-A4B29, A4B33, A4B35, A4B17-A4B23, A4B30-

A4B32, A4B34

(Field Duplicates A4B25/A4B26)

5/Aqueous Equipment Blanks/A4B02, A4B05, A4B06, A4B08,

A4B10

2/Soil PEs/A4B56, A4B57

CERCLIS No. VTD048141741

TDD No. 12-10-0008, Task No. 0850-49

Dear Ms. Bosworth:

A Tier II validation was performed on the organic analytical data for 19 soil samples and five aqueous equipment (rinsate) blanks collected by WESTON START at the Jard Company Inc site in Bennington, Vermont, and for two PE samples obtained from EPA Region I. *Italicized sample ID numbers in the list above are associated with samples in this SDG, but reported in another SDG.* The samples were analyzed under CLP following SOW SOM01.2 as low/medium level for Aroclor compounds. The data were evaluated as Tier II level in accordance with the "Region I EPA-NE Data Validation Functional Guidelines for Evaluating Environmental Analyses" dated December 1996, and the USEPA CLP National Functional Guidelines for Superfund Organic Methods, and were based on the following parameters:

- Overall Evaluation of Data and Potential Usability Issues.
- * Data Completeness.
- * Preservation and Technical Holding Times.
- GC/MS and GC/ECD Instrument Performance Checks.
- * IC and CC.
- * Blanks.
 - Surrogate Compounds.
- NA IS.
- * MS/MSD.

- * Field Duplicates.
- NA Sensitivity Check (MDL Study or LFB).
- * PE Samples/Accuracy Check.
 - Target Compound Identification.
- * Sample Quantitation and Reported Quantitation Limits.
- NA TICs.
- * SVOC and PEST/PCB Cleanup.
- * System Performance.
- NA SEDD/ADR.
- * = No qualifications will be applied based on this parameter.

Table I summarizes overall evaluation of the data with reference to the DQO and potential usability issues. Qualified data are summarized in Data Summary Table 1.

Overall Evaluation of Data and Potential Usability Issues

See Table I for overall evaluation of data and potential usability issues.

Preservation and Technical Holding Times

Aroclor samples A4B17-A4B18, A4B23-A4B29, and A4B30-A4B34 were extracted between 2 and 4 days beyond the holding time specified in SOM01.2. Based upon the holding times articulated in SW-846, the Chlorinated Biphenyl Congener Statement of Work CBC01.2, and in consultation with USEPA Region I Quality Assurance chemists, the holding time for Aroclors has been established as up to 1 year. The positive and non-detected Aroclor results will not be qualified.

Surrogate Compounds

AROCLORS:

Samples in which two or more Aroclor surrogate recoveries did not meet criteria are summarized in the following table:

Sample No.	No. of Surrogates Out	Action Pos/ND
A4B17	4	Accept
A4B17DL	4	Accept
A4B18	4	Accept
A4B18DL	4	Accept
A4B21	4	Accept
A4B21DL	4	Accept

Sample No.	No. of Surrogates Out	Action Pos/ND
A4B22	1	None

Sample results will be qualified as indicated above.

PE Samples/Accuracy Check

The criteria used by START for qualification of sample data based on the PE sample results are as follows:

	Action				
PE Score	Non-Detects	Positive Results			
In Window	Accept	Accept			
Warning Low/High	Accept	Accept			
Action Low	Reject (R)	Estimate (J)			
Action High	Accept	Estimate (J)			
TCL Misses	Reject (R)	Varies			
TCL Contaminants	Accept	Varies			
TIC Misses	Varies	Varies			
TIC Contaminants	Varies	Varies			

All non-compliant PE scores were investigated by checking raw data, calculations, calibrations, possible matrix interferences, and blank contamination. Unless otherwise noted, all results reported by the laboratory were found to be correct, based on the data generated by the laboratory.

The laboratory properly identified and quantified the soil Aroclor-1242 PE sample (A4B56, PE No. ASX0184). No qualifications were applied.

The laboratory properly identified and quantified the soil Aroclor-1260 PE sample (A4B57, PE No. AS1507). No qualifications were applied.

Ms. Martha Bosworth 21 August 2013 Page 4

Target Compound Identification

The dual column correlation did not meet %D confirmation criteria for the following Aroclor compounds:

Sample	Compound	% D	Action
A4B27	Aroclor-1242	72.7	J
A4B23	Aroclor-1242	26.7	J
A4B34	Aroclor-1242	46.4	J

Actions:

J = Estimate results when D > 25 but 00 for pesticides or D > 25 but 00 for PCBs.

R = Reject results when %D >100 for pesticides or %D >500 for PCBs.

U = Qualify result as undetected at the CRQL when %D >100 for pesticides or %D >500 for PCBs, and both results are less than the CRQL.

Sample results have been qualified as indicated above.

Ms. Martha Bosworth 21 August 2013 Page 5

Please contact the undersigned at (978) 552-2100 if you have any questions or need further information.

Very truly yours,

WESTON SOLUTIONS, INC. Region I START

William W. Mahany Principal Project Scientist

Lead Chemist

email cc:

Jennifer Feranda (CLP PO - Region II) - DV Letter w/Data Tables, and ORDA Form only -

Feranda.jennifer@epa.gov

Attachments:

Table I: Overall Evaluation of Soil Data

Data Summary Key Acronym List Data Summary Table 1 DV Worksheets

PE Sample Score Reports (included in DV worksheets)

Field Sampling Notes (including a copy of sampler's COC Records)

CSF Audit (DC-2 Form) - Evidence Audit Photocopy (Including CSF Receipt/Transfer Form)

DQO Summary Form

 $S:\label{lem:case_43392} A4B24\A4B24_val_.doc$

TABLE I

JARD COMPANY INC Case No. 43392; SDG No. A4B24

Overall Evaluation of Soil Data

			AROCLORs		
DQO (list all DQOs)	Sampling and/or			Sampling	
	Analytical Method Appropriate Yes or No	Analytical Error	Sampling Error*	Variability**	Potential Usability Issues
1. To obtain sufficient data from surface and subsurface soil samples collected at the Jard Company site for PCB (Aroclor) analysis, to document potential source areas located on and off the property, and to document contamination in the soil and sediment associated with source areas located on the property.	Analytical Method: Yes, SOM01.2 Sampling Method: Yes, Hand Augers, and Stainless Steel Scoops.	Refer to qualifications in attached Data Summary Table 1.	Refer to qualifications in attached Data Summary Table 1.		1. Positive Aroclor 1242 results in samples A4B23, A4B27, and A4B34 were estimated (J) due to poor dual-column correlation.

^{*} The evaluation of "sampling error" cannot be completely assessed in data validation.

** Sampling variability is not assessed in data validation.

DATA SUMMARY KEY ORGANIC DATA VALIDATION

J The associated numerical value is an estimated quantity. The data are unusable (compound may or may not be present). Resampling and R reanalysis are necessary for verification. The R replaces the numerical value or SQL. The compound was analyzed for, but not detected. The associated numerical IJ value is the SOL or the adjusted SOL. UJ The compound was analyzed for, but not detected. The associated numerical value is the estimated SQL. EB The compound was identified in an aqueous EB that was used to assess field contamination associated with soil/sediment samples. TB The compound was identified in an aqueous TB that was used to assess field contamination associated with soil/sediment samples. BB The compound was identified in an aqueous BB that was used to assess field contamination associated with soil/sediment samples.

ACRONYM LIST ORGANIC DATA VALIDATION

AQ	aqueous	SQL
AQ FB	aqueous field blank	S/S
BB	Bottle Blank	 S/S (m)
B/N	base/neutral compound	START
°C	degrees Celsius	DITHE
CC	-	CVOC
	Continuing Calibration	SVOC
CCV	Continuing Calibration	SW
CT D	Verification	TB
CLP	Contract Laboratory Program	TCL .
COC	Chain-of-Custody record	TDD
COR	Contracting Officer Representative	TIC
CRQL	Contract Required Quantitation	TR
	Limit	VOC
CSF	Complete SDG File	WESTO
%D	percent difference	
DAS	Delivery of Analytical Services	
DMC	Deuterated Monitoring Compound	
DQO	Data Quality Objective	
DV	Data Validation	
DW		
	drinking water	
EB	Equipment Blank	
EPA	Environmental Protection Agency	
GC/ECD	Gas Chromatograph/Electron	
	Capture Detector	
GC/MS	Gas Chromatograph/Mass	
	Spectrometry	
GW	groundwater	
IC	Initial Calibration	
IS	Internal Standard	
kg	kilogram	
L	liter	
LCS	Laboratory Control Sample	
LFB	Laboratory Fortified Blank	
MDL	Method Detection Limit	
	microgram	
μg MS	Matrix Spike	
	- · · · · · · · · · · · · · · · · · · ·	
MSD	Matrix Spike Duplicate	
NA	Not Applicable	
ND	non-detected result	
ng	nanogram	
NERL	New England Regional Laboratory	
OSC	On-Scene Coordinator	
ORDA	Organic Regional Data	
	Assessment	
PAH	polynuclear aromatic hydrocarbon	
PCB	polychlorinated biphenyl	
	compound	
PEST/PCB	pesticide/polychlorinated biphenyl	
1201/102	compound	
PE	Performance Evaluation	
Pos	positive result	
	_	
QC	Quality Control	
%R	percent recovery	
RPD	Relative Percent Difference	
RRF	Relative Response Factor	
RSD	Relative Standard Deviation	
SDG	Sample Delivery Group	
SOW	Statement of Work	
LIDC Defenses #04		

HRS Reference #81

Sample Quantitation Limit soil/sediment soil/sediment medium level Superfund Technical Assessment and Response Team semivolatile organic compound surface water Trip Blank Target Compound List Technical Direction Document Tentatively Identified Compound Traffic Report volatile organic compound ΟN Weston Solutions, Inc.

SITE: JARD COMPANY INC CASE: 43392 SDG: A4B24

LABORATORY: CHEMTECH

CONSULTING GROUP

DATA SUMMARY TABLE 1 AROCLOR IN SOIL ANALYSIS $\mu g/Kg$

	SAMPLE NUMBER SAMPLE LOCATION		A4B24	A4B25	A4B26	A4B27	A4B28	A4B29	A4B33
			SO-07	SO-14	SO-200	SO-21	SO-22	SO-23	SO-62
	STATIC	N LOCATION	JCS-008	JCS-015	JCS-475	JCS-024	JCS-025	JCS-026	JCS-076
	LABORATO	ORY NUMBER	E1902-01	E1902-02	E1902-03	E1902-04	E1902-05	E1902-06	E1902-09
COMPOUND	MDL	CRQL							
Aroclor-1016	2.6	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1221	7.8	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1232	1.3	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1242	6.2	33	41 U	120	130	110 J	140	36 U	180
Aroclor-1248	2.7	33	41 U	37 U	37 U	36 U	- 36 U	36 U	38 U
Aroclor-1254	3.2	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1260	3.2	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1262	14	33	41 U	37 U	37 U	36 U	36 U	36 U	38 U
Aroclor-1268	6.6	33	41 U	- 37 U	37 U	36 U	36 U	36 U	38 U
	DILUT	TION FACTOR	1.0	1.0	1.0	1.0	1.0	1.0	1.0
	DA	TE SAMPLED	4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/3/2013	4/4/2013
	DATE	EXTRACTED	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013
	DAT	E ANALYZED	4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013	4/25/2013
	SAMPLE WEI	GHT (GRAMS)	30	30.1	30.0	30.0	30.0	30.0	30.1
		% SOLID	80.4	88.8	88.4	90.8	91.0	90.6	85.6

NOTES: μg/Kg = micrograms per Kilogram

All results are reported on a Dry Weight Basis.

MDL = Method Detection Limit

CRQL = Contract Required Quantitation Limit

U = Value is Non-Detected.

UJ = Value is Non-Detected, and Detection Limit is Estimated.

J = Value is Estimated.

R = Value is Rejected.

DATA SUMMARY TABLE 1 AROCLOR IN SOIL ANALYSIS $\mu g/Kg$

CONSULTING GROUP

	SAMPLE NUMBER		A4B35	A4B17	A4B18	A4B19	A4B20	A4B21	A4B22
	SAMPLE LOCATION		SO-65	SB-01	SB-03	SB-05	SB-06	SB-08	SB-09
	STATIC	N LOCATION	JCS-086	JCS-130	JCS-136	JCS-138	JCS-148	JCS-153	JCS-143
	LABORATORY NUMBER		E1902-10	E1902-13	E1902-14	E1902-15	E1902-16	E1902-17	E1902-18
COMPOUND	MDL	CRQL							
Aroclor-1016	2.6	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1221	7.8	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1232	1.3	. 33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1242	6.2	33	38 U	280000 *	4800000 *	820 *	1900 *	730000 *	40000 *
Aroclor-1248	2.7	33	38 U	3900 U	3500 U	37 Ú	36 U	3600 U	36 U
Aroclor-1254	3.2	33	38 U	3900 U	3500 U	- 37 U	36 U	3600 U	36 U
Aroclor-1260	3.2	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1262	14	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
Aroclor-1268	6.6	33	38 U	3900 U	3500 U	37 U	36 U	3600 U	36 U
	DILUT	ION FACTOR	1.0	100 / 1000*	100 / 2000*	1 / 2*	1 / 10*	100 / 2000*	1 / 100*
	DA	TE SAMPLED	4/5/2013	4/1/2013	4/1/2013	4/8/2013	4/8/2013	4/8/2013	4/8/2013
DATE EXTRACTED		4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/19/2013	
	DAT	E ANALYZED	4/25/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013	4/26/2013
	SAMPLE WEI	GHT (GRAMS)	30	30	30	30.1	30.1	30.1	30.1
		% SOLID	85.7	84.1	93.5	90.2	90.7	91.0	92.4

NOTES: μg/Kg = micrograms per Kilogram

All results are reported on a Dry Weight Basis.

MDL = Method Detection Limit

CRQL = Contract Required Quantitation Limit

U = Value is Non-Detected.

UJ = Value is Non-Detected, and Detection Limit is Estimated.

J = Value is Estimated.

R = Value is Rejected.

CONSULTING GROUP

DATA SUMMARY TABLE 1 AROCLOR IN SOIL ANALYSIS μg/Kg

	SAMPLE NUMBER		A4B23	A4B30	A4B31	A4B32	A4B34		
	SAMPL	E LOCATION	SO-06	SO-34	SO-36	SO-46	SO-64		
	STATIC	N LOCATION	JCS-006	JCS-046	JCS-048	JCS-061	JCS-183		
	LABORATO	ORY NUMBER	E1902-19	E1902-20	E1902-21	E1902-22	E1902-23		
COMPOUND	MDL	CRQL							
Aroclor-1016	2.6	33	39 U	37 U	39 U	38 U	38 U		
Aroclor-1221	7.8	33	39 U	37 U	39 U	38 U	38 U	:	
Aroclor-1232	1.3	33	39 U	37 U	39 U	· 38 U	38 U		
Aroclor-1242	6.2	. 33	150 J	320	1600 *	1200 *	280 J		
Aroclor-1248	2.7	33	39 U	37 U	39 U	38 U	38 U		
Aroclor-1254	3.2	33	39 U	37 U	39 U	38 U	38 U		
Aroclor-1260	3.2	33	39 U	37 U	39 U	38 U	38 U		
Aroclor-1262	14	33	39 U	37 U	39 U	38 U	38 U		,
Aroclor-1268	6.6	33	39 U	37 U	39 U	38 U	38 U		
				•					
	DILUT	ION FACTOR	1.0	1.0	1 / 10*	1 / 10*	1.0		
	DA	TE SAMPLED	4/3/2013	4/4/2013	4/4/2013	4/4/2013	4/4/2013		
DATE EXTRACTED		4/19/2013	4/19/2013	4/19/2013	4/19/2013	4/22/2013			
	DAT	E ANALYZED	4/25/2013	4/25/2013	4/26/2013	4/26/2013	4/25/2013		
	SAMPLE WEI	GHT (GRAMS)	30	30	30	30	30.1	·	
		% SOLID	85.3	88.6	84.9	88.8	85.6		

NOTES: μg/Kg = micrograms per Kilogram

All results are reported on a Dry Weight Basis.

MDL = Method Detection Limit

CRQL = Contract Required Quantitation Limit

U = Value is Non-Detected.

UJ = Value is Non-Detected, and Detection Limit is Estimated.

J = Value is Estimated.

R = Value is Rejected.

REGION I, EPA-NE ORGANIC REGIONAL DATA ASSESSMENT (ORDA)*

Case No.: 4339 2	Site Name: Jard Company INC
SDG No.: AYBY	No. of Samples/Mat <u>rix: 9 / Soci L</u>
Lab Name: Chemtech Consulting Good	Validation Contracto WESTON
SOW#/Contract#: SOM01.2	Validator's Name: Bill Mahany
EPA-NE DV Tier Level: Tier II	Date DP Rec'd by EPA-NE:
TPO/PO: **ACTION FYI	DV Completion Date: 5/30//3
ANALYTICAL DATA QUALITY S	UMMARY
1. Preservation and Contractual Holding Times: 2. GC/MS / GC/ECD Instrument Performance Check: 3. Initial Calibration: 4. Continuing Calibration: 5. Blanks: 6. DMCs or Surrogate Compounds: 7. Internal Standards: 8. Matrix Spike/Matrix Spike Duplicate: 9. Sensitivity Check: 10. PE samples - Accuracy Check: 11. Target Compound Identification: 12. Compound Quantitation and Reported QLs: 13. Tentatively Identified Compounds: 14. Semivolatile Cleanup/Pesticide/PCB Cleanup: 15. Data Completeness: 16. Overall Evaluation of Data: 0 = Data had no problems or were qualified due to minor contractual m = Data were qualified due to major contractual problems. z = Data were rejected as unusable due to major contractual problem. Action Items (z items):	
Areas of Concern (m items):	
Comments:	
*This form assesses the analytical data quality in items of contractor errors and/or non-contractual analytical issues that affect data quality	
** Check "ACTION" only if contractual defects resulted in reduced Validator:	payment/data rejection recommendations. Date: 5/30//3

Site Name:	Jard Company	Inc
TDD No.:	12-10-0008	
Task No.:	0950	

REGION I ORGANIC DATA VALIDATION

The following data package has been validated:		
Lab Name: Chamtech Consulting Gowp	SOW #/Contract #: SOM	11 2
ن		
Case No.: 4339 2		3, 4, 5, 8, 17
SDG No.: AYBY	Shipping Dates: 4//	7-18/13
No. of Samples/Matrix:	Date Rec'd by Lab:	19-19/13
Traffic Report Sample <u>Nos: AYBJY-B</u>	29 B33, B35, AYBI7 → B	23,830-32,834
Trip Blank No.:		
Equipment Blank No: AY 605, 806, 80	0, Bo 2, B10	
Field Duplicate Nos: <u>AYBYS/BY6</u> PE Nos: AY R56/B57		
PE Nos: <u>AY 856/857</u>		
The <u>Region I, EPA - NE Data Validation Functional Gu</u>		
revision 12/96 was used to evaluate the data and/or ap	·	
Guidelines were used to evaluate the data and are atta		I
criteria from EPA approved QAPjP or amendment to the	ie QAPJP).	
A Tier II or a Tier III evaluation was used to validate the was used, then identify samples, parameters, etc. that	· · · · · · · · · · · · · · · · · · ·	er III
	and the second s	
The data were evaluated based upon the following par	rameters:	
- Overall Evaluation of Data	- Field Duplicates	
- Data Completeness (CSF Audit - Tier I) - Preservation and Technical Holding Times	Sensitivity CheckPE Samples/Accuracy Che	nk
- Freservation and Technical Holding Times - GC/MS and GC/ECD Instrument Performance Check		
- Initial and Continuing Calibrations	- Compound Quantitation and	
- Blanks	Quantitation Limits	
- Surrogate Compounds	-TICs	
- Internal Standards	-Semivolatile and Pesticide/F	CB Cleanup
- Matrix Spike/Matrix Spike Duplicate	- System Performance	·
Region I Definitions and Qualfiers:		
A - Acceptable Data		
J - Numerical value associated with compound is an e	stimated quantity	
R - The data are rejected as unusable. The R replace		n limit
U - Compound not detected at that numerical sample		T minus
UJ - The sample quantitation limit is an estimated qua		
TB, EB - Compound detected in aqueous trip blank or	•	soil/sediment samples.
Rill Mahany		
Validator's Name:	Company Name: <u>WESTON</u> Phone	e Number: <u>978-552-2100</u>
Validator's Name: Rill Mahany Date Validation Started: 5/27/3	Date Validation Completed:	130/13
		-

Data Validation Worksheet Cover Page - Page 2

Check if all criteria are met and no hard copy worksheet provided. Indicate NA if worksheet is not applicable to analytical method. Note: There is no standard worksheet for System Performance, however, the validator must document all system performance issues in the Data Validation Memorandum.

VOA/SV Worksheets:

TABLE II - WORKSHEET OVERALL EVALUATION OF DATA	VOA/SV-Pest/PCB-I PRESI VOA/SV-II GC/MS VOA/SV-III INITIA VOA/SV-IV CONT VOA/SV-Pest/PCB-V-A BLANH VOA/SV-Pest/PCB-V-B BLANH VOA-VI VOA S SV-VI SV SU VOA/SV-VII INTER VOA/SV-Pest/PCB-VIII MATR VOA/SV-Pest/PCB-IX FIELD VOA/SV-Pest/PCB-XI ACCU VOA/SV-Pest/PCB-XII TARG VOA/SV-Pest/PCB-XIII SAMP VOA/SV-XIV TENTA VOA/SV-XV SEMIN	LETE SDG FILE (CSF) AUDIT ERVATION AND HOLDING TIMES INSTRUMENT PERFORMANCE CHECK (TUNING) CALIBRATION NUING CALIBRATION ANALYSIS ANALYSIS URROGATE SPIKE RECOVERIES RROGATE SPIKE RECOVERIES NAL STANDARD PERFORMANCE X SPIKE/MATRIX SPIKE DUPLICATE DUPLICATE PRECISION TIVITY CHECK RACY CHECK/ PE SCORE SHEETS ET COMPOUND IDENTIFICATION LE QUANTITATION ATIVELY IDENTIFIED COMPOUNDS OLATILE CLEANUP ALL EVALUATION OF DATA
---	---	---

Pest/ARO Worksheets:

VOA/SV-Pest/PCB	COMPLETE SDG FILE (CSF) AUDIT	G .
VOA/SV-Pest/PCB-I	PRESERVATION AND HOLDING TIMES	<u> </u>
Pest/PCB-IIA	GC/ECD INSTRUMENT PERFORMANCE CHECK-	
	RESOLUTION	
Pest/PCB-IIB	GC/ECD INSTRUMENT PERFORMANCE CHECK-	
	RETENTION TIMES	
Pest/PCB-IIC	GC/ECD INSTRUMENT PERFORMANCE CHECK-	
	ACCURACY CHECK OF INITIAL CALIBRATION	
Pest/PCB-IID	GC/ECD INSTRUMENT PERFORMANCE CHECK-	
	PESTICIDE DEGRADATION	
Pest/PCB-III	INITIAL CALIBRATION	-
Pest/PCB-IV	CONTINUING CALIBRATION	2-residents
VOA/SV-Pest/PCB-V-A	BLANK ANALYSIS	
VOA/SV-Pest/PCB-V-B	BLANK ANALYSIS	
Pest/PCB-VI	SURROGATE COMPOUNDS: SPIKE RECOVERIES	
	AND RETENTION TIME SHIFT	
Pest/PCB-VII	PESTICIDE CLEANUP	-
VOA/SV-Pest/PCB-VIII	MATRIX SPIKE/MATRIX SPIKE DUPLICATE	
VOA/SV-Pest/PCB-IX	FIELD DUPLICATE PRECISION	
VOA/SV-Pest/PCB-X	SENSITIVITY CHECK	NA
VOA/SV-Pest/PCB-XI	ACCURACY CHECK/ PE SCORE SHEETS	
Pest/PCB-XII	COMPOUND IDENTIFICATION	
VOA/SV-Pest/PCB-XIII	SAMPLE QUANTITATION	
TABLE II - WORKSHEET	OVERALL EVALUATION OF DATA	*

I certify that	t all criteria were j	met for the wo	orksheets checked abov	ve.		@ - always included * - See DV Memo
Signature:	Wmw	2	· · · · · · · · · · · · · · · · · · ·	Name:	Bell	Mahan-1
Date:	5/	113	· · · · · · · · · · · · · · · · · · ·	•		

NA @ NA

COMPLETE SDG FILE (CSF) AUDIT

Organic Fractions:	Aroclor on	14	
Missing Information		Date Lab Contacted	Date Received
		e,	
·			
		·	
			<u> </u>
			·
		en e	
·			·
Validator:M	WMY	Da	te: 5/20/13

ň Of	6. Encore sam	npler (48 hour l
126		Mahar
	Validator:	Mariar

Sampler: G. Wilmin Company: WESTON Contacted: Date:

1. PRESERVATION AND HOLDING TIMES

Cooler 5 6 6 Documented: 747 - 90

Circle sample numbers with exceeded technical holding times or omitted preservation. List all required preservation codes and circle omitted preservation codes.

Circle all exceeded technical holding times.

Identify extraction technique after "# of Days"/(*Extraction Code).

	1					PE	ST					AF	RO		
					# of Days			# of Days			# of Days			# of Days	
Sample No.		Pres.	Date	Date	from Samp.	*Ext.	Date	from Ext.	1	Date	from Samp.	*Ext.	Date	from Ext.	
(TR No.)	Matrix	Code	Sampled	Extracted		Code	Analyzed	to Anal.	Action	Extracted		Code	Analyzed	to Anal.	Action
AYBYY	Soil	. 1	4/3/13	K						4/19/13	16	So X	7/25/13	6	A ¥
BYS.			111			i				1 11/		T,	1 1		11
136															
827															
884	1														
1329											V				
833			7/4/13				- 1				ιŚ				
B35	V	- (415/13				A -				14				
856	PE	1	4/17/13				1 K				ユ		V	V	
657	V		1.7				///				2		4/86/13	7	
AYBIT	Soil		4/1/13								18		Y/26/13		1 *
1318			V							l l	V		1		1
B19			4/8/13							<u> </u>	l t/			(
690															
021										N.				<u> </u>	
372			LV.							4/72/13			W	9	
023			4/3/13							4/19/13	16		Y/25/13	6	4
B 30			4/4/13								15		V	V	
1331			 						Ļ	ļ			4/26/13	. 7	
032	• <i> </i> —		+						1	 		1/	1. 1.	-V	
R34	V		V						<u> </u>	V	V	V	4/25/13	6	VV
			_												
									 						
	L	7	1						<u> </u>	<u> </u>		L			

Preservation Code:

- 1. Cool @ 4°C (± 2°C)
- 2. Preserve with HCl to \leq pH 2.
- 3. Protect from light.
- 4. Freeze.
- 5. Room temperature (avoid excessive heat).
- hold time).

*Extraction Code:

L/L - Liquid/Liquid

SON - Sonication

SEP - Separatory funnel

SOX - Soxhlet

SPE - Solid Phase Extraction

Action Code:

J - Estimate (J) detected values.

UJ - Estimate (UJ) non-detected values.

R - Reject (R) non-detected values.

At = 1-41 holding time let ZIA ar chemist discussions and sw-846/colonis reasons

Matrix Codes:

AQ - Aqueous S/S - Soil/Sediment

AQ FB - Aqueous Field Blank

EPA-NE - Data Validation Worksheet VOA/SV - Pest/ARO - V

V. Rinsate Blank Tabulation - list the applicable rinsate (equipment) blanks below:

Rinsate Blank No.	Sample No.	Equipment Rinsed to Generate the RB	Matrix Applies to:
RB- 0	AY 802	Scol Ro Je	Sail
RB- 02	Bo 5	Hand Auger	
RB- 03	606	1	
RB- 05	1308		
RB- 07	B16	GLOPROBC	V
RB-			

Matrix Codes: SS - surface soil

SD - sediment

SO - source soil

SB - soil boring

GW - groundwater

DW - drinking water

SW - surface water

Note: Apply each RB only to the matrix to which it corresponds. For example, apply the hand auger RB to the soil samples, but not to the surface water samples.

If more than one hand auger/soil sample RB was collected, the RBs may be batched and the highest hit from the batch used to determine the action levels. However, if one RB exhibits an unusual amount of contamination, apply this RB to only the associated samples. Do not batch this RB and apply to all samples of the same matrix.

Validator: Mchan

Date: 5/29/13

EPA - NE -	Data	Validation	Workshee
VOA/SV - F	est/P	CB -XI	

XI. ACCURACY CHECK (Performance Evaluation Results) - List all analytes that are outside criteria.

SDG No.: AYBYY

Case: 43392

Are more than one-half the PE analytes within criteria for each parameter?

Yes

No

Always submit this sheet and attach PE score sheets

PE Sample No.	Ampule No.	Parameter	Type of PE	Matrix	Analyte	Conc.	Region I EPA PE Scores*	Samples Affected	Action
A4056	45×184	50/A60	single blind		Arlaya	UM	PASS	None	Nunc
857	A\$1507	- V	U	V	1360	V	L	10000	77.77
									·
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									,

^{*}For Region I PE indicate the Region I PE Score report result: Action High, Acion Low, TCL Miss, or TCL Contaminant.

Validator:	Mah	any	
	,	1	

Date: 5 (30)/3

PES SCORING EVALUATION REPORT

PES ASX0184

Rev: 2

EPA Sample No.: A4B56

Report Date: 05/13/2013

Page 1 of 1

Lab Name: Chemtech Consulting Group

Contract: EPW11030

Case No.: 43392

Lab Code: CHEM SAS/Client No.: NA

SDG No.: A4B24

Matrix: Soil

Lab Sample ID: E1902-11 Date Extracted: 04/19/2013

Lab File ID: PB004938.D

Date Received: 04/18/2013

% Moisture: 0.0

Date Analyzed: 04/25/2013 Decanted: No

Sample Wt./Vol. (g/mL): 30.0 g

Extraction Type: SOXH

Conc. Extract Vol. (uL): 10000

Injection Vol. (uL): 1.0

GPC Cleanup: No

pH: NA

Sulfur Cleanup: No

Dilution Factor: 1.0

Units: ug/Kg

Analysis Method: SOM01.2 Scoring Method: SOM01.2

Comments:

CAS No.	Analyte	Laboratory Results		PES Evaluation	
		Concentration	Q		
53469-21-9	Aroclor-1242	870	D	PASS	Within Limits
****	END Main Analytes	*****	****	****	*******
****	END All Analytes	******	****	****	******
			-		

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PES SCORING EVALUATION REPORT

PES AS1507

Rev: 1

EPA Sample No.: A4B57

Report Date: 05/13/2013

Page 1 of 1

Lab Name: Chemtech Consulting Group

Contract: EPW11030

Case No.: 43392

Lab Code: CHEM SAS/Client No.: NA

SDG No.: A4B24

Matrix: Soil

Lab Sample ID: E1902-12 Date Extracted: 04/19/2013

Lab File ID: PB004940.D Date Analyzed: 04/26/2013

Date Received: 04/18/2013 Sample Wt./Vol. (g/mL): 30.1 g

% Moisture: 0.0

Decanted: No

Injection Vol. (uL): 1.0

Extraction Type: SOXH

Conc. Extract Vol. (uL): 10000

GPC Cleanup: No

pH: NA

Sulfur Cleanup: No

Dilution Factor: 1.0

Units: ug/Kg

Analysis Method: SOM01.2 Scoring Method: SOM01.2

Comments:

CAS No.	Analyte	Laboratory Results		PES Evaluation		
		Concentration	Q .			
11096-82-5	Aroclor-1260	830	D	PASS	Within Limits	
****	END Main Analytes	*****	****	****	*****	
****	END All Analytes	*****	****	****	******	
	251.05 1111 111111,000					
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EPA - NE - Data Validation Worksheet VOA/SV - Pest/PCB - XIII

XIII. SAMPLE QUANTITATION

If no PE, do sample calculation.

Recalculate, from the raw data, the concentration for one positive detect and one reported sample quantitation limit (SQL) for a non-detect in a diluted sample or soil sample per fraction. (Note: Although Section XIII, C 2. a. requires that one calculation for each fraction in each sample be performed, the validator is only required to reproduce an example, for each fraction, of one positive detect and one SQL calculation on this worksheet.)

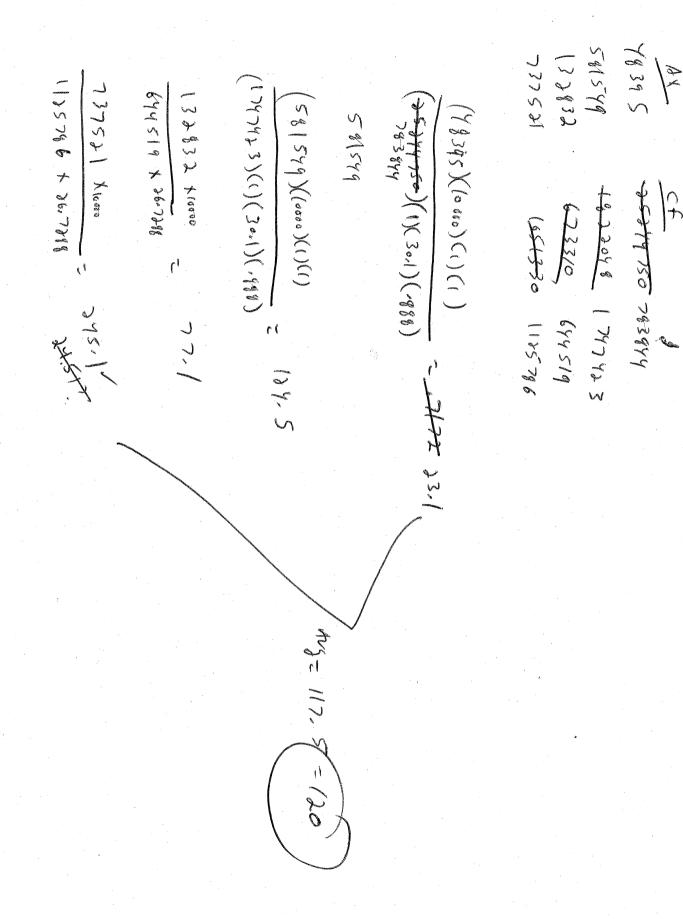
Fraction	Ca	Calculation*				
voc	Detect:	Non-detect QL:				
Sample No.:	The second secon					
Reported Compound:						
Reported Value:						
Non-detected Compound:						
Reported Quantitation Limit:						
svoc	Detect:	Non-detect QL:				
Sample No.:						
Reported Compound:						
Reported Value:						
Non-detected Compound:						
Reported Quantitation Limit:						
P/PCB Sample No.: A4635 Reported Compound: A(1)43- Reported Value:	Detect: $(A_X)(V_T)(0F)(GPC)$ $(F)(V_1)(W_5)(0)$ $(GPC)(GPC)$ $(GPC)(GPC)(GPC)$ $(GPC)(GPC)$ $(GPC)(GPC)(GPC)$ $(GPC)(GPC)(GPC)$ $(GPC)(GPC)(GPC)$ $(GPC)(GPC)$ $(GPC)(G$	Non-detect QL: $33 \times \frac{36}{50.1} \times \frac{1}{858} = 37.038$ = 3 > 0				
Non-detected Compound: M175 Y	- 100 = 0.888	= 370				
Reported Quantitation Limit: 37 U	* MA (T: W/CDE) O/					

* - NA for Tier II if PE score is OK.

Do all soil/sediment samples have % solids greater than 30%? Y N If solids <30%, have sample volumes been increased sufficiently to compensate? Y N If no. list sample numbers_____

Validator: MAM ANY

Date: 5/3°//3



VI. SURROGATE SPIKE RECOVERIES - PESTICIDES/PCBS

List the percent recoveries which do not meet the method QC acceptance criteria.

	TO	CX	DO	CB	Ac	tion
Sample No.	Column 1	Column 2	Column 1	Column 2	Detects	Non-detects
AYBIT	5978	475	376	769	7 A	4
B1 B	19413	530	346	717	(XA	j
Bal	13123	1162	866	868	VA	V
822	378	ව ර්	7.5	85	A	A
AUBITOL	0		0	C	iA	Ps.
AYBISOL	<u> </u>	0	0			
A 40210L	0		<u> </u>	<u> </u>		1
					4	
					13	

		·				
						<u> </u>
			L			

TCX - Tetrachloro-m-xylene

DCB - Decachlorobiphenyl

QC Limits:	30-150	30-150

Qualification of data:

- 1. No action is taken when a sample is analyzed at a dilution.
- 2. No action is required when only one of the four surrogates is outside the QC acceptance criteria and the recovery is > 10%.
- 1. Estimate (J, UJ) all positive and non-detected results if any two surrogates are < the QC acceptance criteria.
- 2. Estimate (J) all positive results if any two surrogates are > the QC acceptance criteria.
- 3. Reject (R) all non-detected results and estimate (J) all positive results if any one surrogate is < 10%.

Sample	One or more	Two or more surrogates	All surrogates	Two or more
Results	surrogates < 10%	10% <u><</u> %R < LL	LL <u>≤</u> %R <u>≤</u> UL	surrogates > UL
Detects	J	J	A	J
Non-detects	R	UJ ·	А	A

LL - Lower Limit

UL - Upper Limit

Validator:	MAWAY	Date:	5/30	13
vanaator.				

100x then 1000 - Zoub

All Criteria Met

Site Name: TARD Company In C

Page ___ of ___

Sample No. A 1 527 A 1 523 A 1 6 3 7	Compound AR 1242 AR 1242 AR 1242	Column I	Column II	%D	Action	Comments
A4627	AR1742	110 150 410	190 190 780	73.7 26.7 46.4	5	
AY 823	AR1242	150	190	76.7	3	
A4834	ARIZYZ	410	780	4604		
		<u> </u>				

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		I	L	L	I	1

Use Comments section to list compounds that went to "U" due to Blank Contamination Actions or Co-elution with Aroclors.

Actions:

- J Estimate results when %D > 25% but \leq 100% for pesticides or %D >25% but \leq 500% for PCBs.
- J@ %D >25% but <100% for pesticides or %D > 25% but <500% for PCBs. Previously qualified as estimated by laboratory due to quantitation below the quantitation limit. No further qualification is needed.
 - R Reject results when %D >100 for pesticides or %D >500% for PCBs.
 - U Qualify result as undetected at the CRQL when %D >100% for pesticides or %D >500% for PCBs and both results are < the CRQL.
- U* Report the non-detected result from the diluted analysis.
- U^ Compound not confirmed by GC/MS. Raise detection limit to reported concentration.
- DL Report the result from the diluted analysis.

	AAA WA
Validator:	MANANY

Date: 5/30//3

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

107, MW-2, MW-3, MW-3D, MW-6, MW-6D, MW-9D, and MW-11. Based on the above information, START personnel planned to purge/develop monitoring wells MW-2, MW-3, MW-3D, MW-6, and MW-6D on 28 March 2013.

1630 hrs: START personnel marked properties located along Park Street and Bowen Road for Dig Safe notification. Following dig safe marking; START personnel secured and departed the site.

28 March 2013 (Thursday) - Site Reconnaissance, Well Development

Weather: Cloudy, high 30 to low 40 °F

0700 hrs: START members Kelly, Hornok, Bitzas, and Robinson arrived at the Jard property. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O₂, H₂S, CO, and PID meter. Background ambient readings: LEL = 0%; O₂ = 20.9%; H₂S = 0 ppm; CO = 0 ppm; and VOC = 0 ppm.

0715 hrs: START HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven terrain, trips-slips-falls, potential weather issues), chemical hazards [PCBs, non-aqueous phase liquids (NAPL) containing water], Radiation (Not encountered previously) and biological hazards (ticks, poison ivy, animals). Personnel reviewed and signed the HASP documentation, as needed.

0800 hrs: START personnel began purging/developing the selected ground water monitoring wells using a Wattera inertia pump system with dedicated tubing, check valve, and surge block at each well. START personnel established on site investigative derived waste (IDW) staging area along west side of Source Pile, on asphalt pavement area/driveway. Location will allow truck for IDW pickup to enter and exit site easily. Staging area consists of 55-gallon drums placed on wooden pallets.

0900 hrs: START PL. Kelly discussed with CORs Bosworth and Smith regarding status of the monitoring well examination, and selection of wells to be purged and sampled. CORs agreed with selection of wells to be sampled.

START personnel continued well purging operations. For the monitoring wells selected for redevelopment/purging, the purge volume in approximate (~) gallons is listed for each well. The following ~ volumes of ground water and/or material were purged from the groundwater wells listed above: MW-2: ~10 gallons; MW-3: ~10 gallons; MW-3D: ~20 gallons; MW-6: ~5 gallons; and MW-6D: ~ 30 gallons. Approximately 4.5 feet of silt material was removed from ground water monitoring well MW-6D. In addition, a very thin NAPL with a greasy feel, along with black oil-like droplets, and a rainbow sheen were observed in IDW purge water

removed from MW-3, MW-3D, and MW-6D.

1330 hrs: START personnel secured the groundwater monitoring well IDW purge water drums, secured the site and departed the Jard property.

1 April 2013 (Monday) - Soil/Source Sampling

Weather: Cloudy, some rain, 45 to 50 °F

1045 hrs: START members Kelly, Hornok, Bitzas, Imbres, Robinson, and Jonathan Saylor arrived at the Jard property.

1100 hrs: START HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven terrain, trips-slips-falls, heavy lifting, Geoprobe Work concerns, potential adverse weather conditions), chemical hazards (PCBs), Radiation (Not encountered previously but will be monitored) and biological hazards (ticks, poison ivy, animals). Personnel reviewed and signed

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

the HASP documentation, as needed. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O_2 , H_2S , CO, and PID meter. Background ambient readings: LEL = 0%; O_2 = 20.9%; O_2 = 0 ppm; O_2 = 0 ppm; and O_2 = 0 ppm. START Team established decontamination area.

1115 hrs: START personnel began decontaminating non-dedicated field sampling equipment including Geoprobe macrocores and cutting shoes, hand augers, metal scoops, and low-flow bladder pumps. Non-dedicated equipment (Geoprobe equipment, augers, metal scoops, etc.) will be decontaminated after the collection of each sample, and prior to use for the collection of other samples.

1400 hrs: Began soil boing activities with the Geoprobe at soil boring location SB-01 located on the south-central area of the former building footprint in an area previously excavated during an EPA Removal action. An EPA removal action was completed at the site during 2007 where the building was razed, a portion of the concrete foundation was removed, and a permeable earthen cap was installed to limit exposure to contaminated soils. Boring activities as part of the Site Reassessment were targeted at the area of the foundation removal and soil excavation (southern portion of the former building footprint).

Sampling on the Jard property and surrounding properties for solid matrices (soil/source, surface soil, and sediment) will be conducted as follows, unless otherwise noted: locations will be designated prior to initiation of sampling activities; at each location, sampling depth will be determined based on sampling objectives and/or materials encountered; for each sampled depth interval at each location, material will be placed in a large polyethylene bag (12 by 15 inches); the material will then be homogenized completely in the bag; the material will later be described by a licensed professional geologist using the modified Burmiester soil classification system and a small sample aliquot will be collected for PCB field screening analysis performed by the US EPA Mobile Laboratory personnel; based on field screening results and sampling objectives, a subset of samples will be selected for further analysis via Contract Laboratory Program (CLP) Aroclor analysis; samples selected for CLP analysis will be aliquoted with sufficient quality assurance/quality control (QA/QC) volume, all solid matrix samples submitted for CLP Aroclor analysis will also be aliquoted for potential congener analysis, unless otherwise noted; following receipt of CLP Aroclor analytical results, a smaller subset of samples will then be selected and submitted for congener analysis. A separate field data sheet will be completed by the field sampler for each sample collected to document relevant information and to supplement field logbook notes.

Additional START personnel performed bump checks on calibrated YSI 550 pH/oxidation reduction potential (ORP)/Conductivity probes for ground water sampling scheduled to be completed on 2 April 2013. All the calibrated ground water sampling equipment was working properly (See calibration log sheets).

1415 hrs: Soil/source sample SB-01A (Sample #: JCS-128) was collected using a Geoprobe macrocore from a depth of 2.7 to 4 feet bgs from soil boring SB-01 and later submitted for PCB field screening analysis.

1420 hrs: Soil/source sample SB-01B (Sample #: JCS-129) was collected using a Geoprobe macrocore from a depth of 6.9 to 8 feet bgs from soil boring SB-01 and later submitted for PCB field screening analysis.

1430 hrs: Soil/source sample SB-01C (Sample #: JCS-130) was collected using a Geoprobe macrocore from a depth of 10.4 to 12 feet bgs from soil boring SB-01 and later submitted for PCB field screening analysis.

1440 hrs: Soil/source sample SB-01D (Sample #: JCS-131) was collected using a Geoprobe macrocore from a depth of 12 to 14 feet bgs from soil boring SB-01 and later submitted for PCB field screening analysis.

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 1500 hrs: START personnel completed soil boing activities at location SB-01. Soil boring SB-01 was completed to a depth of 14 feet bgs due to refusal. See the soil Boring Logs for complete descriptions of the boring completed. The soil boring was backfilled with sand and bentonite. START personnel relocated to and began boring activities at soil boring location SB-02 located on the south-central area of the former building footprint in an area previously excavated during an EPA Removal action.
- 1520 hrs: Soil/source sample SB-02A (Sample #: JCS-132) was collected using a Geoprobe macrocore from a depth of 2.2 to 4 feet bgs from soil boring SB-02 and later submitted for PCB field screening analysis.
- 1530 hrs: Soil/source sample SB-02B (Sample #: JCS-133) was collected using a Geoprobe macrocore from a depth of 6.9 to 8 feet bgs from soil boring SB-02 and later submitted for PCB field screening analysis.
- 1540 hrs: Soil/source sample SB-02C (Sample #: JCS-134) was collected using a Geoprobe macrocore from a depth of 8.8 to 10 feet bgs from soil boring SB-02 and later submitted for PCB field screening analysis.
- 1545 hrs: START personnel completed soil boing activities at location SB-02. Soil boring SB-02 was completed to a depth of 10 feet bgs due to refusal. Evidence (piece of) the orange snow fence layer installed as part of the earthen cap construction was encountered at 2.5 feet bgs. See the soil Boring Logs for complete descriptions of the boring completed. The soil boring was backfilled with sand and bentonite. START personnel relocated to and began boring activities at soil boring location SB-03, located on the south-central capped area, adjacent to ground water monitoring wells MW-3 and MW-3D.
- 1555 hrs: Soil/source sample SB-03A (Sample #: JCS-135) was collected using a Geoprobe macrocore from a depth of 0.7 to 2.6 feet bgs from soil boring SB-03 and later submitted for PCB field screening analysis.
- 1605 hrs: Soil/source sample SB-03B (Sample #: JCS-136) was collected using a Geoprobe macrocore from a depth of 4.8 to 6.5 feet bgs from soil boring SB-03 and later submitted for PCB field screening analysis.
- 1610 hrs: START personnel completed soil boing activities at location SB-03. Soil boring SB-03 was completed to a depth of 6.5 feet bgs due to refusal. The soil boring was backfilled with sand and bentonite. START personnel completed soil boring activities for the day.
- 1630 hrs: Equipment rinsate blank sample RB-01 (Sample #: JCW-013; CLP #: A4B02) was collected from the Geoprobe macrocore system sampling equipment and is associated with soil/source sampling activities conducted on 1 April 2013.
- 1700 hrs: START personnel secured the site and departed the Jard property.

2 April 2013 (Tuesday) - Ground Water Sampling

Weather: Cloudy, little precipitation, low 30 °F

- 0700 hrs: START members Kelly, Hornok, Bitzas, Imbres, Robinson, and Saylor arrived at the Jard property.
- 0715 hrs: START HSC Kelly HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven terrain, trips-slips-falls, potential weather issues), chemical hazards [PCBs, non-aqueous phase liquids (NAPL) containing water], Radiation (Not encountered previously) and biological hazards (ticks, poison ivy, animals). Personnel reviewed and signed the HASP documentation, as needed. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O₂, H₂S, CO, and PID meter. Background ambient readings: LEL = 0%; O₂ = 20.9%; H₂S = 0 ppm; CO = 0 ppm; and VOC = 0 ppm. Note that the ground

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

water monitoring equipment was calibrated on 1 April 2013 and bump-tested on 2 April 2013, and determined to be with calibration specifications (see calibration sheets).

START Team established decontamination area and conduct decontamination of non-dedicated equipment. Non-dedicated equipment (bladder pumps, measuring tapes, etc.) will be decontaminated after the collection of each sample, and prior to use for the collection of other samples.

0830 hrs: START member Bitzas began monitoring of low-flow parameters at ground water monitoring well EPA-104D located in the wetland area, west of Park Street and downgradient from the Jard property, behind the residential properties. START member Bitzas monitored low-flow ground water parameters per the Site-Specific Quality Assurance project Plan (QAPP) and START standard operating procedures (SOPs). See the field data sheets for more information.

0905 hrs: START member Saylor began monitoring of low-flow parameters at ground water monitoring well MW-11 located northwest of the Jard property. START member Saylor monitored low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information.

0920 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-10 [Matrix Spike/Matrix Spike Duplicate (MS/MSD)] (Sample #: JCW-010; CLP #: A4A99) was collected from monitoring well EPA-104D. Ground water sample GW-10 was collected using a bladder pump and the final water quality parameters were as follows: Temperature = 3.74 degrees Celsius (°C); Specific Conductivity = 91 micro Siemens per centimeter (μS/cm); pH = 5.72; ORP = 229.3 millivolts (mv); Dissolved Oxygen (DO) = 9.11 milligrams per liter (mg/L); and turbidity = 21.2 Nephelometric Turbidity Units (NTU). Note: Due to a YSI 550 probe malfunction, an additional volume of sample was collected for pH and ORP measurements using a second YSI 550. In addition, pH was monitored during low-flow activities with pH paper. Readings on pH paper indicated a pH between 5.0 and 6.0. A total of approximately 13.5 liters was purged prior to sample collection with the pump intake at 20.0 ft below the TOC. See the field data sheets for more information.

0940 hrs: START member Imbres began monitoring of low-flow parameters at ground water monitoring well EPA-107 located northwest of the Jard property. START member Imbres monitored low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information.

1000 hrs: START member Robinson began monitoring of low-flow parameters at ground water monitoring well MW-9D located west of the Jard property. START member Robinson monitored low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information.

1030 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-02 (Sample #: JCW-002; CLP #: A4A91) was collected from monitoring well EPA-107. Ground water sample GW-02 was collected using a bladder pump and the final water quality parameters were as follows: Temperature = 5.14 °C; Specific Conductivity = 162 μ S/cm; pH = 7.12; ORP = 192.9 mv; DO = 4.40 mg/L; and turbidity = 0.51 NTU. A total of approximately 11.0 liters was purged prior to sample collection with the pump intake at 17 ft. below the TOC.

11.0 liters was purged prior to sample collection with the pump intake at 17 ft. below the 10C.

1110 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-09
(Sample #: JCW-009; CLP #: A4A98) was collected from monitoring well MW-11. Ground water sample GW-09 was collected using a bladder pump and the final water quality parameters were as follows: Temperature = 3.42 °C; Specific Conductivity = 55 µS/cm; pH = 6.34; ORP = 138.6 mv; DO = 11.41 mg/L; and turbidity = 10.3 NTU. A total of approximately 61.7 liters was purged prior to sample collection with the pump intake at 6.5 ft. below the TOC. The turbidity meter initially used to evaluate low-flow ground water parameters compliance malfunctioned and was replaced with one that was operational.

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 1125 hrs: Stabilization of water quality parameters was achieved and groundwater sample GW-08 (Sample #: JCW-008; CLP #: A4A97) and field duplicate GW-11 (Sample #: JCW-011; CLP #: A4B00) were collected from monitoring well MW-9D located west of the Jard property. Ground water samples GW-08 and GW-11 were collected using a bladder pump and the final water quality parameters were as follows: Temperature = 6.52 °C; Specific Conductivity = 44 µS/cm; pH = 6.29; ORP = 25.2 mv; DO = 0.13 mg/L; and turbidity = 34.2 NTU. A total of approximately 17 liters was purged prior to sample collection with the pump intake at 24 ft below the TOC. An issue was encountered with the turbidity meter during low-flow ground water monitoring, and was replaced with one that was operating.
- 1200 hrs: pH and ORP measurements were collected from a volume of water from ground water sample GW-10 and were as follows: pH = 5.72; ORP = 229.3 mv.
- 1250 hrs: START member Imbres began monitoring of low-flow parameters at ground water monitoring well EPA-100 located north of the Jard property. START member Imbres monitored all low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information.
- 1310 hrs: START member Bitzas began monitoring of low-flow parameters at ground water monitoring well MW-2 located on the southern portion of the Jard property. START member Bitzas monitored all low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information.
- 1345 hrs: START member Saylor began monitoring of low-flow parameters at ground water monitoring well MW-3D located directly south of the former building footprint on the southern portion of the Jard property. START member Saylor monitored all low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information.
- 1355 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-01 (Sample #: JCW-001; CLP #: A4A90) was collected from monitoring well EPA-100. Ground water sample GW-01 was collected using a bladder pump and the final water quality parameters were as follows: Temperature = 4.03 °C; Specific Conductivity = 119 μS/cm; pH = 6.61; ORP = 266.3 mv; DO = 11.74 mg/L; and turbidity = 1.78 NTU. A total of approximately 13.8 liters was purged prior to sample collection with the pump intake at 32 ft below the TOC.
- 1345 hrs: START member Robinson began monitoring of low-flow parameters at ground water monitoring well MW-6D located directly west of the former building footprint on the western boundary of the Jard property. START member Robinson monitored all low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information.
- 1405 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-03 (Sample #: JCW-003; CLP #: A4A92) was collected from monitoring well MW-02. Ground water sample GW-03 was collected using a peristaltic pump with dedicated tubing and the final water quality parameters were as follows: Temperature = 1.41 °C; Specific Conductivity = 79 μ S/cm; pH = 5.59; ORP = 175.3 mv; DO = 11.41 mg/L; and turbidity = 0.91 NTU. A total of approximately 11 liters was purged prior to sample collection with the intake at 8.6 ft below the TOC.
- 1505 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-05 (Sample #: JCW-005; CLP #: A4A94) was collected from monitoring well MW-3D. Ground water sample GW-05 was collected using a bladder pump and the final water quality parameters were as follows: Temperature = 1.90 °C; Specific Conductivity = 47 μS/cm; pH = 6.37; ORP = 112.7 mv; DO = 4.75 mg/L; and turbidity = 1.16 NTU. A total of approximately 13.6 liters was purged prior to sample collection with the pump intake at 29 ft below the TOC. In addition, stabilization of water quality parameters was achieved and ground water sample GW-07 (Sample #: JCW-007; CLP # A4A96) was collected from monitoring well MW-6D.

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

Ground water sample GW-07 was collected using a bladder pump and the final water quality parameters were as follows: Temperature = 7.17 °C; Specific Conductivity = 42 μ S/cm; pH = 6.53; ORP = 203.9 mv; DO = 8.80 mg/L; and turbidity = 51.7 NTU. A total of approximately 16.7 liters was purged prior to sample collection with the pump intake at 26.5 ft below the TOC.

1545 hrs: START member Robinson began monitoring of low-flow parameters at ground water monitoring well MW-6 located directly west of the former building footprint on the western boundary of the Jard property. START member Robinson monitored all low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information. The YSI initially used to monitor low-flow ground water parameters was replaced with one that was operational.

1555 hrs: START member Saylor began monitoring of low-flow parameters at ground water monitoring well MW-3 located directly south of the former building footprint on the southern portion of the Jard property. START member Saylor monitored all low-flow ground water parameters per the Site-specific QAPP and START SOPs. See the field data sheets for more information. Low-flow ground water parameters were not conducted within a flow cell due to potential contamination/non-aqueous phase liquid (NAPL) within the well. Previous purging of the well on 28 March 2013 indicated product within the well that had a greasy feel and contained small oil droplets that were black in color.

1600 hrs: Equipment rinsate blank sample RB-20 (Sample #: JCW-012; CLP #: A4B01) was collected from a bladder pump sampling equipment and is associated with ground water sampling activities.

1630 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-06 (Sample #: JCW-006; CLP #: A4A95) was collected from monitoring well MW-6. Ground water sample GW-06 was collected using a peristaltic pump with dedicated tubing and the final water quality parameters were as follows: Temperature = 4.18 °C; Specific Conductivity = 116 μS/cm; pH = 6.35; ORP = -83.6 mv; DO = 0.20 mg/L; and turbidity = 0.72 NTU. A total of approximately 9 liters was purged prior to sample collection with the intake at 13.5 ft below the TOC.

1700 hrs: Stabilization of water quality parameters was achieved and ground water sample GW-04 (Sample #: JCW-004; A4A93) was collected from monitoring well MW-3. Ground water sample GW-04 was collected using a peristaltic pump with dedicated tubing and the final water quality parameters were as follows: Temperature = 1.25 °C; Specific Conductivity = 69 μS/cm; pH = 6.67; ORP = -158.2 mv; DO = 4.40 mg/L; and turbidity = 0.93 NTU. A total of approximately 15 liters was purged prior to sample collection with the intake at 10.5 ft below the TOC. Ground water sample GW-04 was also collected for congener analysis.

1730 hrs: IDW purge water was containerized in 55-gallon steel drums and segregated based on well location (on or off the Jard property). Segregation is to aid in later IDW disposal, assuming wells from on the Jard property contain high concentrations of contaminants than those wells located off the Jard property. In addition, waste soil material and IDW Decon waste are also segregated to aid in later IDW disposal activities. START personnel secured IDW drums, secured the site and departed the Jard property.

3 April 2013 (Wednesday) – Soil/Source Sampling

Weather: Partly cloudy, high 30 °F

0700 hrs: START members Kelly, Hornok, Bitzas, Imbres, Robinson, and Jonathan Saylor arrived at the Jard property. In addition, performance evaluation samples PE-AA3325 (Sample #: JCW-014;

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

CLP #: A4B03) and PE-AA2555 (Sample #: JCW-015; CLP#: A4B04) were collected for CLP Aroclor analysis.

- 0715 hrs: START HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven terrain, trips-slips-falls, heavy lifting, potential adverse weather conditions), chemical hazards (PCBs), Radiation (Not encountered previously but will be monitored) and biological hazards (ticks, poison ivy, snakes, animals). Personnel reviewed and signed the HASP documentation, as needed. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O₂, H₂S, CO, and PID meter. Background ambient readings: LEL = 0%; O₂ = 20.9%; H₂S = 0 ppm; CO = 0 ppm; and VOC = 0 ppm.
 - START Team established decontamination area and conduct decontamination of non-dedicated equipment. Non-dedicated equipment (augers, metal scoops, etc.) will be decontaminated after the collection of each sample, and prior to use for the collection of other samples.
- 0815 hrs: Soil/source sample SO-01A (Sample #: JCS-001) was collected with a hand auger at a depth of 0 to 8 inches bgs from the upper north-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 0820 hrs: Soil/source sample SO-02A (Sample #: JCS-002) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 0830 hrs: Soil/source sample SO-03A (Sample #: JCS-003) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper north-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-04A (Sample #: JCS-004) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 0840 hrs: Soil/source sample SO-05A (Sample #: JCS-005) was collected with a hand auger at a depth of 0 to 8 inches bgs from the upper north-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-06A (Sample #: JCS-006) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 0845 hrs: Soil/source sample SO-06B (Sample #: JCS-007) and soil/source field duplicate SO-100B (Sample #: JCS-065) were collected with a hand auger at a depth of 6 to 12 inches bgs from the upper central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 0855 hrs: Soil/source sample SO-07A (Sample #: JCS-008) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper north-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis. In addition, soil/source sample SO-08A (Sample #: JCS-009) was collected with a hand auger at a depth of 0 to 8 inches bgs from the upper north-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1010 hrs: Soil/source sample SO-09A (Sample #: JCS-027) was collected with a hand auger at a depth of 0 to 8 inches bgs from the upper west-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-10A (Sample #: JCS-010) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

- 1025 hrs: Soil/source sample SO-12A (Sample #: JCS-013) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper east-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1030 hrs: Soil/source sample SO-11A (Sample #: JCS-011) was collected with a hand auger at a depth of 0 to 18 inches bgs from the upper west-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1035 hrs: Soil/source sample SO-11B (Sample #: JCS-012) was collected with a hand auger at a depth of 18 to 42 inches bgs from the upper west-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis,
- 1040 hrs: Soil/source sample SO-14A (Sample #: JCS-015) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper east-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1050 hrs: Soil/source sample SO-16A (Sample #: JCS-017) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper south-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1055 hrs: Soil/source sample SO-16B (Sample #: JCS-018) was collected with a hand auger at a depth of 6 to 12 inches bgs from the upper south-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1100 hrs: Soil/source sample SO-13A (Sample #: JCS-014) was collected with a hand auger at a depth of 0 to 16 inches bgs from the upper western-central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1106 hrs: Soil/source sample SO-18A (Sample #: JCS-020) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper southwestern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1110 hrs: Soil/source sample SO-15A (Sample #: JCS-016) was collected with a hand auger at a depth of 0 to 18 inches bgs from the upper southwestern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1123 hrs: Soil/source sample SO-18B (Sample #: JCS-021) was collected with a hand auger at a depth of 6 to 12 inches bgs from the upper southwestern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1125 hrs: Soil/source sample SO-17A (Sample #: JCS-019) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper southwestern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1130 hrs: Soil/source sample SO-19A (Sample #: JCS-022) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper central portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1143 hrs: Soil/source sample SO-20A (Sample #: JCS-023) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper southern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1150 hrs: Soil/source sample SO-21A (Sample #: JCS-024) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper southern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1155 hrs: Soil/source sample SO-22A (Sample #: JCS-025) was collected with a hand auger at a depth of 0 to 6 inches bgs from the upper southwestern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- Soil/source sample SO-23A (Sample #: JCS-026) was collected with a hand auger at a depth of 1200 hrs: 0 to 3 inches bgs from the upper southwestern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

Jard Company, Inc.

Bennington, Vermont

CERCLIS No. VTD048141741

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 1215 hrs: START personnel continued to complete CLP documentation and to package ground water and rinsate blank samples for shipment to the CLP Laboratory located in Mountainside, New Jersey. START geologist Kelly continued to conduct classification of sample matrix materials using the modified Burmiester soil classification and to prepare sample aliquots for field screening.
- 1540 hrs: Equipment rinsate blank sample RB-02 (Sample #: JCW-016: CLP #: A4B05) was collected from a hand auger sampling equipment (augers, scoops, etc.) and is associated with soil/source sampling activities.
- 1630 hrs: START personnel completed sample shipment preparation, organized and packaged traffic reports. START member Bitzas left the site and proceeded to deliverer samples and paperwork to FedEx, located in Menands, New York for shipment. Below is a summary of the traffic reports (TR), Airbill numbers (AB), and samples sent to the CLP Organics Laboratory (Chemtech Consulting Group) for PCB Aroclor analysis:
 - TR #: 1-040313-081601-0001, Master AB #: 5141 2418 0581, four groundwater samples for PCB Aroclor analysis. These four samples were shipped as dangerous goods due to previous sampling results and field observations and were to be combined with samples shipped under TR #: 1-040313-083108-0002 AB #: 5141 2418 0559, to constitute a complete sample delivery group (SDG) with appropriate quality assurance/quality control (QA/QC) samples.
 - TR #: 1-040313-083108-0002 AB #: 5141 2418 0559, seven ground water samples including one field duplicate, and one MS/MSD; plus two rinsate blank, and two performance evaluation samples for PCB Aroclor analysis. Samples from this TR were to be combined with samples shipped under TR #: 1-040313-081601-0001, Master AB #: 5141 2418 0581, to form a complete SDG.

1700 hrs: START personnel secured IDW drums, secured the site and departed the Jard property.

4 April 2013 (Thursday) - Soil/Source Sampling

Weather: Sunny, 45 to 50 °F

- 0700 hrs: START members Kelly, Hornok, Bitzas, Imbres, Robinson, and Jonathan Saylor arrived at the Jard property.
- 0715 hrs: START HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven terrain, trips-slips-falls, heavy lifting, traffic concerns, potential adverse weather conditions), chemical hazards (PCBs), Radiation (Not encountered previously but will be monitored) and biological hazards (ticks, poison ivy, animals). Personnel reviewed and signed the HASP documentation, as needed. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O2, H2S, CO, and PID meter. Background ambient readings: LEL = 0%; O2 = 20.9%; H2S = 0 ppm; CO = 0 ppm; and VOC = 0 ppm.
 - START Team established decontamination area and conduct decontamination of non-dedicated equipment. Non-dedicated equipment (augers, metal scoops, etc.) will be decontaminated after the collection of each sample, and prior to use for the collection of other samples.
- 0800 hrs: Soil/source sample SO-24A (Sample #: JCS-078) was collected with a hand auger at a depth of 0 to 8 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.
 - In addition, soil/source sample SO-25A (Sample #: JCS-028) was collected with a hand auger at a depth of 0 to 12 inches bgs from the drainage ditch located on the western portion of the Jard property and later submitted for PCB field screening analysis.

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- Soil/source sample SO-50A (Sample #: JCS-066) was collected with a hand auger at a depth of 0 to 12 inches bgs from the area below the former transformer area located on the southern portion of the Jard property and later submitted for PCB field screening analysis.
- 0805 hrs: Soil/source sample SO-24B (Sample #: JCS-079) was collected with a hand auger at a depth of 8 to 24 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.
- 0810 hrs: Soil/source sample SO-25B (Sample #: JCS-029) was collected with a hand auger at a depth of 12 to 30 inches bgs from the drainage ditch located on the western portion of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-24C (Sample #: JCS-080) was collected with a hand auger at a depth of 24 to 30 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.

Soil/source sample SO-50B (Sample #: JCS-067) was collected with a hand auger at a depth of 12 to 16 inches bgs from the area below the former transformer area located on the southern portion of the Jard property and later submitted for PCB field screening analysis.

- 0815 hrs: Soil/source sample SO-25C (Sample #: JCS-030) was collected with a hand auger at a depth of 30 to 48 inches bgs from the drainage ditch located on the western portion of the Jard property and later submitted for PCB field screening analysis.
- 0818 hrs: Soil/source sample SO-51A (Sample #: JCS-068) was collected with a hand auger at a depth of 0 to 6 inches bgs from the area below the former transformer area located on the southern portion of the Jard property and later submitted for PCB field screening analysis.
- 0823 hrs: Soil/source sample SO-26A (Sample #: JCS-031) was collected with a hand auger at a depth of 0 to 12 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.
- 0826 hrs: Soil/source sample SO-52A (Sample #: JCS-069) was collected with a hand auger at a depth of 0 to 4 inches bgs from the area below the former transformer area located on the southern portion of the Jard property and later submitted for PCB field screening analysis.
- 0830 hrs: Soil/source sample SO-27A (Sample #: JCS-036) was collected with a hand auger at a depth of 0 to 18 inches bgs from the drainage ditch located on the western portion of the Jard property and later submitted for PCB field screening analysis.
- 0833 hrs: Soil/source sample SO-26B (Sample #: JCS-032) was collected with a hand auger at a depth of 12 to 18 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.
- 0835 hrs: Soil/source sample SO-27B (Sample #: JCS-038) was collected with a hand auger at a depth of 18 to 24 inches bgs from the drainage ditch located on the western portion of the Jard property and later submitted for PCB field screening analysis.
- 0836 hrs: Soil/source sample SO-26C (Sample #: JCS-033) was collected with a hand auger at a depth of 18 to 24 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.
- 0840 hrs: Soil/source sample SO-26D (Sample #: JCS-034) was collected with a hand auger at a depth of 24 to 36 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.
- 0845 hrs: Soil/source sample SO-29A (Sample #: JCS-040) was collected with a hand auger at a depth of . 0 to 12 inches bgs from the area located along the western boundary of the Jard property and later submitted for PCB field screening analysis.
- 0850 hrs: Soil/source sample SO-26E (Sample #: JCS-035) was collected with a hand auger at a depth of 36 to 42 inches bgs from the drainage ditch located on the northwestern portion of the Jard property and later submitted for PCB field screening analysis.

Jard Company, Inc.

Bennington, Vermont

CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 0900 hrs: Soil/source sample SO-31A (Sample #: JCS-043) was collected with a hand auger at a depth of 0 to 12 inches bgs from the area located along the western boundary of the Jard property and later submitted for PCB field screening analysis.
- 0905 hrs: Soil/source sample SO-28A (Sample #: JCS-039) was collected with a hand auger at a depth of 0 to 8 inches bgs from the area located along the western boundary of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-31B (Sample #: JCS-044) was collected with a hand auger at a depth of 12 to 24 inches bgs from the area located along the western boundary of the Jard property and later submitted for PCB field screening analysis.
- 0920 hrs: Soil/source sample SO-30A (Sample #: JCS-041) was collected with a hand auger at a depth of 0 to 12 inches bgs from the area located along the southwestern boundary of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-33A (Sample #: JCS-081) was collected with a hand auger at a depth of 0 to 18 inches bgs from the area located along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
- 0930 hrs: Soil/source sample SO-30B (Sample #: JCS-042) was collected with a hand auger at a depth of 12 to 24 inches bgs from the area located along the southwestern boundary of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-33B (Sample #: JCS-082) was collected with a hand auger at a depth of 18 to 30 inches bgs from the area located along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
- 0935 hrs: Soil/source sample SO-33C (Sample #: JCS-083) was collected with a hand auger at a depth of 30 to 36 inches bgs from the area located along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
- 0940 hrs: Soil/source sample SO-32A (Sample #: JCS-045) was collected with a hand auger at a depth of 0 to 12 inches bgs from the area located along the southwestern boundary of the Jard property and later submitted for PCB field screening analysis.
- 1035 hrs: Soil/source sample SO-35A (Sample #: JCS-047) was collected with a hand auger at a depth of 0 to 12 inches bgs from the southwestern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1045 hrs: Soil/source sample SO-37A (Sample #: JCS-049) was collected with a hand auger at a depth of 0 to 6 inches bgs from the southwestern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-53A (Sample #: JCS-084) was collected with a hand auger at a depth of 0 to 12 inches bgs from the area below the former transformer area located on the southern portion of the Jard property and later submitted for PCB field screening analysis.
- 1055 hrs: Soil/source sample SO-39A (Sample #: JCS-051) was collected with a hand auger at a depth of 0 to 12 inches bgs from the southwestern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-54A (Sample #: JCS-085) was collected with a hand auger at a depth of 0 to 8 inches bgs from the area below the former transformer area located on the southern portion of the Jard property and later submitted for PCB field screening analysis.
- 1100 hrs: Soil/source sample SO-39B (Sample #: JCS-052) was collected with a hand auger at a depth of 12 to 24 inches bgs from the southwestern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 START geologist Kelly continued to conduct classification of sample matrix materials using the modified Burmiester soil classification and to prepare sample aliquots for field screening.

- 1110 hrs: Soil/source sample SO-41A (Sample #: JCS-054) was collected with a hand auger at a depth of 0 to 8 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1120 hrs: Soil/source sample SO-41B (Sample #: JCS-055) was collected with a hand auger at a depth of 8 to 18 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1125 hrs: Soil/source sample SO-41C (Sample #: JCS-056) was collected with a hand auger at a depth of 18 to 30 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1145 hrs: Soil/source sample SO-34A (Sample #: JCS-046) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper northeastern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis. In addition, soil/source sample SO-43A (Sample #: JCS-058) was collected with a hand auger at a depth of 0 to 12 inches bgs from the southwestern toe slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1150 hrs: Soil/source sample SO-45A (Sample #: JCS-060) was collected with a hand auger at a depth of 0 to 18 inches bgs from the western toe slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1210 hrs: Soil/source sample SO-47A (Sample #: JCS-062) was collected with a hand auger at a depth of 0 to 6 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1210 hrs: START Member Hornok contacted and discussed sampling progress with COR Bosworth. Discussed number of samples collected to date, groundwater well sampling status, difficulties source sampling to depth on the upper portion of the source pile, source areas along western property boundary, and planned field screening and sampling activities. Scott Clifford (EPA Chemist) will be on site on Monday (4/8/13).
- 1225 hrs: Soil/source sample SO-38A (Sample #: JCS-050) was collected with a hand auger at a depth of 0 to 8 inches bgs from the upper northern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1230 hrs: Soil/source sample SO-36A (Sample #: JCS-048) was collected with a hand auger at a depth of 0 to 12 inches bgs from the upper northern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-49A (Sample #: JCS-064) was collected with a plastic scoop at a depth of 0 to 3 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1235 hrs: Soil/source sample SO-55A (Sample #: JCS-070) was collected with a plastic scoop at a depth of 0 to 4 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1350 hrs: Soil/source sample SO-56A (Sample #: JCS-071) was collected with a hand auger at a depth of 0 to 12 inches bgs from the western toe slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1405 hrs: Soil/source sample SO-57A (Sample #: JCS-072) was collected with a hand auger at a depth of 0 to 6 inches from the western toe slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1415 hrs: Soil/source sample SO-40A (Sample #: JCS-053) was collected with a hand auger at a depth of 0 to 8 inches bgs from the upper northwestern portion of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

Jard Company, Inc. Bennington, Vermont CERCLIS No. VTD048141741

DRAFT Log Book Notes

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 1420 hrs: Soil/source sample SO-42A (Sample #: JCS-057) was collected with a hand auger at a depth of 0 to 12 inches bgs from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1425 hrs: Soil/source sample SO-59A (Sample #: JCS-074) was collected with a metal scoop at a depth of 0 to 4 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1430 hrs: Soil/source sample SO-58A (Sample #: JCS-073) was collected with a metal scoop at a depth of 0 to 2 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1435 hrs: Soil/source sample SO-60A (Sample #: JCS-075) was collected with a hand auger at a depth of 0 to 12 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1438 hrs: Soil/source sample SO-46A (Sample #: JCS-061) was collected with a hand auger at a depth of 0 to 8 inches from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1440 hrs: Soil/source sample SO-44A (Sample #: JCS-059) was collected with a hand auger at a depth of 0 to 6 inches bgs from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1505 hrs: Soil/source sample SO-61A (Sample #: JCS-182) was collected with a hand auger at a depth of 0 to 12 inches from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1515 hrs: Soil/source sample SO-63A (Sample #: JCS-077) was collected with a hand auger at a depth of 0 to 8 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1520 hrs: Soil/source sample SO-62A (Sample #: JCS-076) was collected with a hand auger at a depth of 0 to 12 inches from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-64A (Sample #: JCS-183) was collected with a hand auger at a depth of 0 to 4 inches from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1630 hrs: Soil/source sample SO-48A (Sample #: JCS-063) was collected with a plastic scoop at a depth of 0 to 3 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1700 hrs: Equipment rinsate blank sample RB-03 (Sample #: JCW-017: CLP #: A4B06) was collected from hand auger sampling equipment (augers, scoops, etc.) associated with soil/source sampling activities.
- 1705 hrs: Equipment rinsate blank sample RB-04 (Sample #: JCW-018: CLP #: A4B07) was collected from hand auger sampling equipment (augers, scoops, etc.) associated with soil/source sampling activities.
- 1730 hrs: START personnel secured IDW drums, secured the site and departed the Jard property.

5 April 2013 (Friday) – Soil/Source Sampling

Weather: Partly cloudy, low 50 °F

- 0730 hrs: START members Kelly, Hornok, Bitzas, Imbres, Robinson, and Jonathan Saylor arrived at the Jard property. COR Bosworth also arrived on site for meeting with EPA and town representatives.
- 0745 hrs: START HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

terrain, trips-slips-falls, heavy lifting, traffic concerns, potential adverse weather conditions), chemical hazards (PCBs), Radiation (Not encountered previously but will be monitored) and biological hazards (ticks, poison ivy, animals). Personnel reviewed and signed the HASP documentation, as needed. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O2, H2S, CO, and PID meter. Background ambient readings: LEL = 0%; $O_2 = 20.9\%$; $O_2 = 0$ ppm; $O_3 = 0$ ppm; and $O_3 = 0$ ppm.

START Team established decontamination area and conduct decontamination of non-dedicated equipment. Non-dedicated equipment (augers, metal scoops, etc.) will be decontaminated after the collection of each sample, and prior to use for the collection of other samples.

START embers Kelly and Hornok held discussions with COR Bosworth regarding current status of sampling activities, Flex-viewer Data Management Resource, and groundwater shipment/delivery.

0845 hrs: Soil/source sample SO-65A (Sample #: JCS-086) was collected with a hand auger at a depth of 0 to 8 inches from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

START geologist Kelly continued to conduct classification of sample matrix materials using the modified Burmiester soil classification and to prepare sample aliquots for field screening.

0855 hrs: Soil/source sample SO-66A (Sample #: JCS-087) was collected with a plastic scoop at a depth of 0 to 3 inches bgs from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

0900 hrs: Soil/source sample SO-67A (Sample #: JCS-088) was collected with a hand auger at a depth of 0 to 6 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-68A (Sample #: JCS-089) was collected with a hand auger at a depth of 0 to 12 inches bgs from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

0915 hrs: Soil/source sample SO-70A (Sample #: JCS-093) was collected with a hand auger at a depth of 0 to 12 inches bgs from the northern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-72A (Sample #: JCS-095) was collected with a hand auger at a depth of 0 to 10 inches bgs from the drainage area at the base of the northeastern corner of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

0920 hrs: Soil/source sample SO-69A (Sample #: JCS-090) was collected with a hand auger at a depth of 0 to 12 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-72B (Sample #: JCS-096) was collected with a hand auger at a depth of 10 to 20 inches bgs from the drainage area at the base of the northeastern corner of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

0925 hrs: Soil/source sample SO-69B (Sample #: JCS-091) was collected with a hand auger at a depth of 12 to 36 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

0930 hrs: Soil/source sample SO-69C (Sample #: JCS-092) was collected with a hand auger at a depth of 36 to 48 inches bgs from the western slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 0945 hrs: Soil/source sample SO-76A (Sample #: JCS-102) was collected with a hand auger at a depth of 0 to 14 inches bgs from the eastern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 0950 hrs: Soil/source sample SO-71A (Sample #: JCS-094) was collected with a hand auger at a depth of 0 to 24 inches bgs from the lower eastern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-74A (Sample #: JCS-098) was collected with a hand auger at a depth of 0 to 12 inches bgs from the eastern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1000 hrs: Soil/source sample SO-74B (Sample #: JCS-099) was collected with a hand auger at a depth of 12 to 30 inches bgs from the eastern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1005 hrs: Soil/source sample SO-73A (Sample #: JCS-097) was collected with a hand auger at a depth of 0 to 28 inches bgs from the eastern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1010 hrs: Soil/source sample SO-75A (Sample #: JCS-100) was collected with a hand auger at a depth of 0 to 12 inches bgs from the eastern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1020 hrs: Soil/source sample SO-77A (Sample #: JCS-101) was collected with a hand auger at a depth of 0 to 18 inches bgs from the eastern slope of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.
- 1100 hrs: VT DEC ANR Wetland Specialist Julie Foley arrived on-site to discuss wetland areas around the site with START personnel. Wetland Specialist Foley provide START with previously completed wetland delineation map of wetlands to west of Park Street. START members Hornok, Bitzas and Kelly reviewed available wetland references/maps with Specialist Foley.
- 1130 hrs: Equipment rinsate blank sample RB-05 (Sample #: JCW-019: CLP #: A4B08) was collected from hand auger sampling equipment (augers, scoops, etc.) associated with soil/source sampling activities.
 - COR Bosworth returned from meeting with Section Chief Meghan Cassidy to review operations.
- 1140 hrs: COR Bosworth and Section Chief Cassidy departed site.

 Wetland Specialist Foley accompanied START members Kelly and Bitzas on reconnaissance of wetland areas to the west of Park Street and background wetland area along Bowen Road north of the Jard property.
- 1200 hrs: START personnel completed sample shipment preparation, organized and packaged traffic reports. START members Robinson and Saylor proceeded to deliverer samples and paperwork to FedEx, located in Brattleboro, VT for shipment. Below is a summary of the TRs AB numbers and samples sent to the CLP Organics Laboratory (Chemtech Consulting Group) for PCB Aroclor analysis:
 - TR #: 1-040513-111321-0003, Master AB #: 5141 2418 0662, four aqueous equipment rinsate blank samples for PCB Aroclor analysis.
- 1215 hrs: Wetland Specialist Foley confirmed that wetland delineation map of wetlands to west of Park Street is generally the same as current conditions based on reconnaissance and review of area. Foley also noted that the proposed background area to the north along Bowen Road, contained similar types of wetland (PEM, PSS, POW, etc.). Wetland Specialist Foley left the site to attend a local meeting/inspection.
 - START member Kelly spoke START PM McDuffee regarding leaving the Geoprobe Truck secured on the VTrans property for the weekend in an effort to be more sustainable/"Green". PM McDuffee agreed to plan.

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

START members Hornok and Kelly spoke with a VTrans representative at the Bowen Road facility regarding leaving the Geoprobe Truck secured on the VTDOT property for the weekend in an effort to be more sustainable/"Green". The VTrans representative agreed and explained there security for the weekend hours and where it would be best to park the vehicle. Informed PM McDuffee agreed to plan.

1230 hrs: Remaining START personnel secured IDW drums, secured the site and departed the Jard property for the START office located in Andover, MA.

8 April 2013 (Monday) – Soil/Source Sampling

Weather: Partly sunny, high 50 to low 60 °F

- 1030 hrs: START members Kelly, Hornok, Bitzas, Eric Ackerman, Chris Dupree, Robinson, Jonathan Saylor, and Robert Sharp arrived at the Jard property. START Member Hornok picked up Geoprobe truck from VT DOT facility along Bowen Road. In addition, EPA SAM Martha Bosworth had already arrived on-site.
- 1045 hrs: START HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven terrain, trips-slips-falls, heavy lifting, Geoprobe Work concerns, potential adverse weather conditions), chemical hazards (PCBs), Radiation (Not encountered previously but will be monitored) and biological hazards (ticks, poison ivy, dogs, animals). Personnel reviewed and signed the HASP documentation, as needed. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O2, H2S, CO, and PID meter. Background ambient readings: LEL = 0%; O2 = 20.9%; H2S = 0 ppm; CO = 0 ppm; and VOC = 0 ppm.

START Team established decontamination area and conduct decontamination of non-dedicated equipment. Non-dedicated equipment (Geoprobe equipment, augers, metal scoops, etc.) will be decontaminated after the collection of each sample, and prior to use for the collection of other samples.

- 1100 hrs: Soil boring activities began at soil boring location SB-05 located on the south-eastern area of the former building footprint in an area previously excavated during an EPA Removal action. In addition, EPA Office of Environmental Measurement and Evaluation (OEME) Mobile Laboratory chemist Scott Clifford arrived on-site to perform PCB field screening analysis. Sample aliquots for PCB field screening, collected to date between 1 April and 5 April, were transferred to EPA chemist Clifford for processing and PCB field screening analyses. START geologist Kelly continued to conduct classification of sample matrix materials using
- the modified Burmiester soil classification and to prepare sample aliquots for field screening.

 1130 hrs: Soil/source sample SB-05A (Sample #: JCS-137) was collected using a Geoprobe macrocore from a depth of 2.1 to 4 feet bgs from soil boring SB-05 and later submitted for PCB field screening analysis.
- 1135 hrs: Soil/source sample SB-05B (Sample #: JCS-138) was collected using a Geoprobe macrocore from a depth of 5.3 to 5.6 feet bgs from soil boring SB-05 and later submitted for PCB field screening analysis.
- 1140 hrs: START personnel completed soil boring activities at location SB-05. Soil boring SB-05 was completed to a depth of 6 feet bgs due to refusal. Team backfilled hole with sand and bentonite and relocated to next location. Boring activities began at soil boring location SB-07 located on the south-eastern area of the former building footprint in an area previously excavated during an EPA Removal action.

PCB field screening analysis.

DRAFT Log Book Notes

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 1145 hrs: Soil/source sample SB-07A (Sample #: JCS-139) was collected using a Geoprobe macrocore from a depth of 2 to 2.9 feet bgs from soil boring SB-07 and later submitted for PCB field screening analysis.
- 1155 hrs: START personnel completed soil boring activities at location SB-07. Soil boring SB-07 was completed to a depth of 4 feet bgs due to refusal. Team backfilled hole with sand and bentonite and relocated to next location. Boring activities began at soil boring location SB-09 located on the south-eastern area of the former building footprint in an area previously excavated during an EPA Removal action. In addition, boring activities began at soil boring location SB-04 located beneath the former transformer area located on the southern portion of the Jard property.
- 1210 hrs: Soil/source sample SB-09A (Sample #: JCS-140) was collected using a Geoprobe macrocore from a depth of 2.9 to 3.4 feet bgs from soil boring SB-09 and later submitted for PCB field screening analysis.
 - In addition, soil/source sample SB-09B (Sample #: JCS-141) was collected using a Geoprobe macrocore from a depth of 3.4 to 4 feet bgs from soil boring SB-09 and later submitted for PCB field screening analysis.
- 1215 hrs: START Member Kelly decided to collect an additional sample from upper core section to obtain analyses throughout the core section. Soil/source sample SB-09C (Sample #: JCS-142) was collected using a Geoprobe macrocore from a depth of 1.7 to 2.9 feet bgs from soil boring SB-09 and later submitted for PCB field screening analysis.
- 1220 hrs: Soil/source sample SB-04A (Sample #: JCS-145) was collected using a Geoprobe macrocore from a depth of 1.1 to 1.3 feet bgs from soil boring SB-04 and later submitted for PCB field screening analysis.

 In addition, soil/source sample SB-04B (Sample #: JCS-146) was collected using a Geoprobe macrocore from a depth of 1.3 to 2 feet bgs from soil boring SB-04 and later submitted for
- 1230 hrs: START personnel completed soil boring activities at location SB-04. Soil boring SB-04 was completed to a depth of 2 feet bgs due to refusal. Team backfilled hole with sand and bentonite and relocated to next location. Boring activities began at soil boring location SB-06 located on the south-western area of the former building footprint in an area previously excavated during an EPA Removal action.
- 1235 hrs: Soil/source sample SB-06A (Sample #: JCS-147) was collected using a Geoprobe macrocore from a depth of 2.3 to 3.3 feet bgs from soil boring SB-06 and later submitted for PCB field screening analysis.

 In addition, soil/source sample SB-06B (Sample #: JCS-148) was collected using a Geoprobe

macrocore from a depth of 3.3 to 4 feet bgs from soil boring SB-06 and later submitted for PCB field screening analysis.

- 1240 hrs: After reviewing the entire core, START Member Kelly decided to collect an additional sample from upper core section to obtain analyses throughout the core to represent various depths. Soil/source sample SB-06C (Sample #: JCS-149) was collected using a Geoprobe macrocore from a depth of 1.5 to 2.3 feet bgs from soil boring SB-06 and later submitted for PCB field screening analysis.
- 1245 hrs: START personnel completed soil boring activities at location SB-06. Soil boring SB-06 was completed to a depth of 4 feet bgs due to refusal. Team backfilled hole with sand and bentonite and relocated to next location.
- 1250 hrs: Soil/source sample SB-09D (Sample #: JCS-143) was collected using a Geoprobe macrocore from a depth of 7.4 to 8 feet bgs from soil boring SB-09 and later submitted for PCB field screening analysis.

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

- 1255 hrs: Soil/source sample SB-09E (Sample #: JCS-144) was collected using a Geoprobe macrocore from a depth of 10.1 to 11 feet bgs from soil boring SB-09 and later submitted for PCB field screening analysis.
- 1300 hrs: START personnel completed soil boring activities at location SB-09. Soil boring SB-09 was completed to a depth of 11 feet bgs. Team backfilled hole with sand and bentonite and relocated to next location. Boring activities began at soil boring location SB-08 located on the south-eastern area of the former building footprint in an area previously excavated during an EPA Removal action.
- 1345 hrs: Soil/source sample SB-08A (Sample #: JCS-150) was collected using a Geoprobe macrocore from a depth of 1.2 to 4 feet bgs from soil boring SB-08 and later submitted for PCB field screening analysis.
- 1350 hrs: Soil/source sample SB-08B (Sample #: JCS-151) was collected using a Geoprobe macrocore from a depth of 6.9 to 8 feet bgs from soil boring SB-08 and later submitted for PCB field screening analysis.
- 1400 hrs: Soil/source sample SB-08C (Sample #: JCS-152) was collected using a Geoprobe macrocore from a depth of 8.7 to 10 feet bgs from soil boring SB-08 and later submitted for PCB field screening analysis.

 In addition, soil/source sample SB 08D (Sample #: JCS 153) was collected using a Geoprobe

In addition, soil/source sample SB-08D (Sample #: JCS-153) was collected using a Geoprobe macrocore from a depth of 10 to 11 feet bgs from soil boring SB-08 and later submitted for PCB field screening analysis.

- 1410 hrs: START personnel completed soil boring activities at location SB-08. Soil boring SB-08 was completed to a depth of 11 feet bgs due to equipment issues (stuck). Team did not backfilled hole, will work to retrieve equipment later and backfill with sand and bentonite; relocated to next location. Boring activities began at soil boring location SB-10 located on the eastern edge of the former building footprint in an area previously excavated during an EPA Removal action.
 - In addition, soil/source sample SO-81A (Sample #: JCS-106) was collected with a hand auger at a depth of 0 to 18 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
- 1415 hrs: Soil/source sample SO-80A (Sample #: JCS-103) was collected with a hand auger at a depth of 0 to 18 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
 - In addition, soil/source sample SO-82A (Sample #: JCS-109) was collected with a hand auger at a depth of 0 to 18 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
 - 1420 hrs: Soil/source sample SO-81B (Sample #: JCS-107) was collected with a hand auger at a depth of 18 to 36 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
 - 1425 hrs: Soil/source sample SO-80B (Sample #: JCS-104) was collected with a hand auger at a depth of 18 to 30 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-81C (Sample #: JCS-108) was collected with a hand auger at a depth of 36 to 54 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-82B (Sample #: JCS-110) was collected with a hand auger at a depth of 18 to 30 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

- 1435 hrs: Soil/source sample SO-80C (Sample #: JCS-105) was collected with a hand auger at a depth of 30 to 40 inches bgs from an area along the northwestern boundary of the Jard property and later submitted for PCB field screening analysis.
- 1445 hrs: Soil/source sample SO-83A (Sample #: JCS-111) was collected with a hand auger at a depth of 0 to 12 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.

 In addition, soil/source sample SO-84A (Sample #: JCS-112) was collected with a hand auger.

In addition, soil/source sample SO-84A (Sample #: JCS-112) was collected with a hand auger at a depth of 0 to 18 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.

- 1450 hrs: Soil/source sample SO-85A (Sample #: JCS-114) was collected with a hand auger at a depth of 0 to 12 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.
- 1455 hrs: Soil/source sample SO-84B (Sample #: JCS-113) and soil/source sample field duplicate SO-102B (Sample #: JCS-207) were collected with a hand auger at a depth of 18 to 36 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.
- 1500 hrs: Soil/source sample SO-85B (Sample #: JCS-115) and soil/source sample field duplicate SO-101B (Sample #: JCS-206) were collected with a hand auger at a depth of 12 to 24 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.
- 1510 hrs: Soil/source sample SO-85C (Sample #: JCS-116) was collected with a hand auger at a depth of 24 to 30 inches from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-86A (Sample #: JCS-117) was collected with a hand auger at a depth of 0 to 18 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.

Soil/source sample SO-87A (Sample #: JCS-118) was collected with a hand auger at a depth of 0 to 18 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.

- 1520 hrs: Soil/source sample SO-87B (Sample #: JCS-119) was collected with a hand auger at a depth of 18 to 36 inches bgs from an area along the northern boundary of the Jard property and later submitted for PCB field screening analysis.
- 1530 hrs: Soil/source sample SO-88A (Sample #: JCS-120) was collected with a hand auger at a depth of 0 to 18 inches bgs from an area along the eastern edge of the building footprint on the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-89A (Sample #: JCS-122) was collected with a hand auger at a depth of 0 to 12 inches bgs from the northeastern corner of the Jard property and later submitted for PCB field screening analysis.

- 1535 hrs: Soil/source sample SO-88B (Sample #: JCS-121) was collected with a hand auger at a depth of 18 to 30 inches bgs from along the eastern edge of the building footprint on the Jard property and later submitted for PCB field screening analysis.
- 1540 hrs: Soil/source sample SO-89B (Sample #: JCS-123) was collected with a hand auger at a depth of 12 to 24 inches bgs from the northeastern corner of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SO-90A (Sample #: JCS-124) was collected with a hand auger at a depth of 0 to 18 inches bgs from along the eastern edge of the pile located on the eastern portion of the Jard property and later submitted for PCB field screening analysis.

Jard Company, Inc.
Bennington, Vermont
CERCLIS No. VTD048141741

TDD No. 01-12-10-0008 Logbook No. 01433-S Site Reassessment

1545 hrs: Soil/source sample SO-91A (Sample #: JCS-125) was collected with a hand auger at a depth of 0 to 10 inches from the northeastern corner of the Jard property and later submitted for PCB field screening analysis.

In addition, soil/source sample SB-10A (Sample #: JCS-154) was collected using a Geoprobe macrocore from a depth of 0.4 to 1.3 feet bgs from soil boring SB-10 and later submitted for PCB field screening analysis.

1600 hrs: START personnel completed soil boring activities at location SB-10. Soil boring SB-10 was completed to a depth of 2 feet bgs due to refusal. Team backfilled sample hole with sand and bentonite.

Soil/source sample SO-92A (Sample #: JCS-126) was collected with a hand auger at a depth of 0 to 8 inches from the northeastern corner of the Jard property and later submitted for PCB field screening analysis.

- 1615 hrs: Soil/source sample SO-93A (Sample #: JCS-127) was collected with a plastic scoop at a depth of 0 to 2 inches bgs from an area along the northeastern edge of the building footprint on the Jard property and later submitted for PCB field screening analysis.
- 1635 hrs: Equipment rinsate blank sample RB-06 (Sample #: JCW-020: CLP #: A4B09) was collected from hand auger sampling equipment (augers, scoops, etc.) associated with soil/source sampling activities.
- 1640 hrs: Equipment rinsate blank sample RB-07 (Sample #: JCW-021: CLP #: A4B10) was collected from the Geoprobe macrocore system sampling equipment and is associated with soil/source sampling activities.
- 1700 hrs: START personnel secured IDW drums, secured the site and departed the Jard property.

9 April 2013 (Tuesday) – Soil/Source and Surface Soil Sampling

Weather: Cloudy, high 50 to low 60 °F

- 0700 hrs: START members Kelly, Hornok, Bitzas, Ackerman, Dupree, Robinson, Saylor, and Sharp arrived at the Jard property. EPA SAM Martha Bosworth had previously arrived on-site. In addition, Chemist Clifford also arrived on-site.
- 0715 hrs: START HSC Kelly reviewed the site HASP and conducted a tailgate health and safety meeting for all on-site START personnel, including reviews of the physical hazards (uneven terrain, trips-slips-falls, heavy lifting, traffic, potential adverse weather conditions), chemical hazards (PCBs), Radiation (Not encountered previously but will be monitored) and biological hazards (ticks, poison ivy, animals). Personnel reviewed and signed the HASP documentation, as needed. START members completed calibration checks on air monitoring instrument; MultiRAE Plus, LEL, O2, H2S, CO, and PID meter. Background ambient readings: LEL = 0%; O2 = 20.9%; H2S = 0 ppm; CO = 0 ppm; and VOC = 0 ppm.

START Team established decontamination area and conduct decontamination of non-dedicated equipment. Non-dedicated equipment (augers, metal scoops, etc.) will be decontaminated after the collection of each sample, and prior to use for the collection of other samples.

- 0800 hrs: START members Kelly, Robinson, and Scesny began marking sample locations and documenting property features on the Park Street residential properties.
- 0810 hrs: Soil/source sample SO-95A (Sample #: JCS-185) was collected with a hand auger at a depth of 0 to 8 inches bgs from an area along the northwestern edge of the building footprint on the Jard property and later submitted for PCB field screening analysis.
- 0815 hrs: Soil/source sample SO-94A (Sample #: JCS-184) was collected with a hand auger at a depth of 0 to 12 inches bgs from an area along the northwestern edge of the building footprint on the Jard property and later submitted for PCB field screening analysis.

Page 1 of 1

USEPA CLP Organics COC (FAB COPY)

DateShipped: 4/18/2013 CarrierName: FedEx AirbillNo: 5141 2418 0938 **CHAIN OF CUSTODY RECORD**

COPY

No: 1-041713-114538-0006

Lab: ChemTech Consulting Group Lab Contact: Divya Mehta Lab Phone: 908-789-8900

SDA# AU

Case #: 43392 Cooler #: DG Cans

Organic Sample #	Matrix/Sampler	Coll. Method	Analysis/Turnaround	Tag/Preservative/Bottles	Station Location	Collected	Inorganic Sample #	For Lab Use Only
A4B17	Soil/ START	Grab	CLP PCBs(21)	537 (4 C) (1)	JCS-130	04/01/2013 14:30		
A4B18	Soil/ START	Grab	CLP PCBs(21)	538 (4 C) (1)	JCS-136	04/01/2013 16:05		
A4B19	Soil/ START	Grab	CLP PCBs(21)	539 (4 C) (1)	JCS-138	04/08/2013 11:35		
A4B20	Soil/ START	Grab	CLP PCBs(21)	. 540 (4 C) (1)	JCS-148	04/08/2013 12:35		
A4B21	Soil/ START	Grab	CLP PCBs(21)	541 (4 C) (1)	JCS-153	04/08/2013 14:00		
A4B22	Soil/ START	Grab	CLP PCBs(21)	542 (4 C) (1)	JCS-143	04/08/2013 12:50		
A4B23	Soil/ START	Grab	CLP PCBs(21)	543 (4 C) (1)	JCS-006	04/03/2013 08:40		200
A4B30	Soil/ START	Grab	CLP PCBs(21)	565 (4 C) (1)	JCS-046	04/04/2013 11:45		
A4B31 ·	Soil/ START	Grab	CLP PCBs(21)	567 (4 C) (1)	JCS-048	04/04/2013 12:30		
A4B32 ·	Soil/ START	Grab	CLP PCBs(21)	569 (4 C) (1)	JCS-061	04/04/2013 14:38		
A4B34	Soil/ START	Grab	CLP PCBs(21)	573 (4 C) (1)	JCS-183	04/04/2013 15:20		
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Special Instructions: Samples shipped as dangerous goods due to elevated PCB concentrations noted in field screening analysis. Please combine these samples with samples shipped under Fedex Airbill # 5141 2418 0743, COC #: 1-041713-115310-0007 to create one SDG.

Shipment for Case Complete? N

Samples Transferred From Chain of Custody #

Analysis Key: CLP PCBs=SOM01.2 Aroclors

Items/Reason	Relinquished by	Date	Received by	Date	Time	Items/Reason	Relinquished By	Date	Received by	Date	Time
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AUB34 is least sample of this sDa #

Mage 55 of 126

Page 1 of 1

USEPA CLP Organics COC (LAB COPY)

DateShipped: 4/17/2013 CarrierName: FedEx AirbillNo: 5141 2418 0743



CHAIN OF CUSTODY RECORD

Case #: 43392 Cooler #: MM1276 No: 1-041713-115310-0007

Lab: ChemTech Consulting Group Lab Contact: Divya Mehta Lab Phone: 908-789-8900

Organic Sample #	Matrix/Sampler	Coll. Method	Analysis/Turnaround	Tag/Preservative/Bottles	Station Location	Collected	Inorganic Sample #	For Lab Use Only
A4B24	Soil/ START	Grab	CLP PCBs(21)	544 (4 C) (1)	JCS-008	04/03/2013 08:55		
A4B25	Soil/ START	Grab	CLP PCBs(21)	553 (4 C) (1)	JCS-015	04/03/2013 10:40		
A4B26	Soil/ START	Grab	CLP PCBs(21)	556 (4 C) (1)	JCS-475	04/03/2013 10:40		
A4B27	Soil/ START	Grab	CLP PCBs(21)	558 (4 C) (1)	JCS-024	04/03/2013 11:50		
A4B28	Soil/ START	Grab	CLP PCBs(21)	560 (4 C) (1)	JCS-025	04/03/2013 11:55		
A4B29	Soil/ START	Grab	CLP PCBs(21), CLP PCBs(21)	562 (4 C), 563 (4 C) (2)	JCS-026	04/03/2013 12:00		
A4B33	Soil/ START	Grab	CLP PCBs(21)	571 (4 C) (1)	JCS-076	04/04/2013 15:20		
A4B35	Soil/ START	Grab	CLP PCBs(21)	575 (4 C) (1)	JCS-086	04/05/2013 08:45		
A4B56 ? E	Soil/ START	Grab	CLP PCBs(21)	620 (4 C) (1)	JCS-478	04/17/2013 07:00		
A4B57 P7	Soil/ START	Grab	CLP PCBs(21)	621 (4 C) (1)	JCS-479	04/17/2013 07:00		
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Sample(s) to be used for Lab QC: A4B29 - Special Instructions: Please combine with samples shipped under FedEx Airbill #: 5141 2418 0710, COC #: 1-041713-114538-0006 to form one sample delivery group.

Shipment for Case Complete? N
Samples Transferred From Chain of Custody #

Analysis Key: CLP PCBs=SOM01.2 Aroclors

Items/Reason	Relinquished by	Date	Received by	Date	Time	Items/Reason	Relinquished By	Date	Received by	Date	Time
Sampus	C forman	4(1713	ALTON VO.	4/17/13	1700						
										and the second s	
										t.	
						Souphis	2141 2418 0313		Gorge VEGRO.	14)18/13	945
								- O.C -		,	

Temp: 500

Page 56 of 126 / (

DG/ESAY

2.
 3.
 4.

LABORATORY NAME:

ORGANICS COMPLETE SDG FILE (CSF) INVENTORY SHEET FORM DC-2

CHEMTECH CONSULTING GROUP, INC.

Jard Company Weston

CITY / STATE :	МО	UNTAINSIDE, NJ		U		
CASE NO:	43392	SDG NO : A4B24		C. C	MAY 0 9	2013
SDG NOs TO FOL	LOW	N/A N/A			7	
MOD. REF. NO. :		N/A		N/A B	1	
CONTRACT NO:		EPW11030				
SOW NO:		SOM 01.2			MAY 09	2043
All documents deliv	ered in t	he Complete SDG File (CSF) must be original doc	uments where p	ossible.		
			PA FROM	GE NOS: TO	CH LAB	USEPA/Weshi
Inventory Sheet	(DC-2) (Do not number)		10		
SDG Narrative			1	8	~	
SDG Cover Shee	et/Traffic	Report	9	11		
Trace Volatiles I				•		
a. QC Summa	-	toring Compound Recovery (Form II VOA-1				
and VOA		compound Necovery (Form II VOA-1	NA	NA		
		rix Spike Duplicate Recover (Form III VOA) JSEPA Region)	NA	NA		
Method E	Blank Su	mmary (Form IV VOA)	NA	NA		·
GC/MS I	nstrumer	nt Performance Check (Form V VOA)	NA	NA		
Internal S	Standard	Area and RT Summary (Form VIII VOA)	, NA	NA		
b. Sample Dat TCL Res and VOA	ults - Or	ganics Analysis Data Sheet (Form I VOA-1	NA	NA		
Tentative	ly Identii	fied Compounds (Form I VOA-TIC)				
Reconstr	ucted to	tal ion chromatograms (RIC) for each sample				
For each	sample): :				
		and background-subtracted mass spectra of unds identified			Market State Control of the Control	
Quant	titation re	eports				
		of all reported TICs with three best library matche	s NA	NA		
Initial Ca	libration	Data (Form VI VOA-1, VOA-2, VOA-3)				
RICs and	d Quantit	taation Reports for all Standards				
Continuir	ng Calibr	ration Data (Form VII VOA-1, VOA-2, VOA-3)				
RICs and	l Quantil	tation Reports for all Standards				
d. Raw/Quality	Control		•			
BFB			NA	NA	1	
Blank Data	a		NA	NA		
Matrix Spil USEPA Re		x Spike Duplicate Data (if requested by	NA	NA		V
	•	Evidence Audit Photo	copy			

CASE	NO: 43392	SDG NO: A4B24	SDG NOs	TO FOLLOW	. N/A	**************************************	
N/A		N/A	MOD. REF	. NO :	N/A		
е.	Trace SIM Data (Pla	ace at the end of the Trace V	olatiles Section	NA	NA		し
	IV-VOA-SIM; For	l; Form II VOA-SIM1 and VOATM VI VOA-SIM; Form VII VOATM All raw data for QC, Sample	A-SIM; Form				
Low	v/Med Volatiles Data						
a.	QC Summary						
	Deuterated MonivOA-2, VOA-3, \	toring Compound Recovery (I/OA-4)	Form II VOA-1,	NA	<u>NA</u>		
		rix Spike Duplicate Recovery equested by USEPA Region)	(Form III VOA-1	NA	NA NA		
	Method Blank Su	ımmary (Form IV VOA)		NA:	<u>NA</u>		
	GC/MS Instrume	nt Performance Check (Form	V VOA)	NA	<u>NA</u>		
	Internal Standard	d Area and RT Summary (For	m VIII VOA)	NA	NA		
b.	Sample Data			NA	NA	_	
	TCL Results - Or VOA-1 and VOA	ganics Analysis Data Sheet (-2)	Form I				
	Tentatively Identi	fied Compounds (Form I VO	A-TIC)				
	Reconstructed to	tal ion chromatograms (RIC)	for each sample				
		e: a and background-subtracted ounds identified	mass spectra of				
	Quantitation	reports					*
	Mass Spectr matches	a of all reported TICs with thr	ee best library				
c.	Standards Data (Al	l Instruments		NA	NA		. 1
	Initial Calibration	Data (Form VI VOA-1, VOA-	2, VOA-3)				
	RICs and Quanti	tation Reports for all Standard	ds				W .
	Continuing Calib	ration Data (Form VII VOA-1,	VOA-2, VOA-3)				
	RICs and Quanti	tation Reports for all Standar	ds				
d.	Raw/Quality Contro	ol (QC)Data					
	BFB			NA .	NA		
	Blank Data			NA	NA		
	Matrix Spike/Mat USEPA Region)	rix Spike Duplicate Data (if re	quested by	NA	NA		V

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CASE NO: 43392	SDG NO: A4B24	SDG NOs TO FOLLOW:	N/A
N/A	N/A	MOD. REF. NO: N/A	

6. <u>Semivolatiles Data</u>

a.	QC Summary				
	Deuterated Monitoring Compound Recovery (Form II SV-1, SV-2, SV-3, SV-4)	NA	NA		4
	Matrix Spike/Matrix Spike Duplicate Recovery Summary (Form III SV-1 and SV-2) (if requested by USEPA Region)	NA	NA		
	Method Blank Summary (Form IV SV)	NA	NA		
	GC/MS Instrument Performance Check (Form V SV)	NA	NA		
	Internal Standard Area and RT Summary (Form VIII SV-1 and SV-2)	NA	<u>NA</u>		
b.	Sample Data	NA	NA	•	
	TCL Results - Organics Analysis Data Sheet (Form I SV-1 and SV-2)				
	Tentatively Identified Compounds (Form I SV-TIC)		•		
	Reconstructed total ion chromatograms (RIC) for each sample				
	For each sample:	NA	NA	· _	
	Raw Spectra and background-subtracted mass spectra of target compounds				
	Quantitation reports				
	Mass Spectra of TICs with three best library matches			minute of the contract of the	
	GPC chromatograms (if GPC is r				
c.	Standards Data (All Instruments)	NA	NA	_	
	Initial Calibration Data (Form VI SV-1, SV-2, SV-3)				
	RICs and Quantitation				
	Continuing Calibration Data (Form VII SV-1, S				
	RICs and Quantitation Reports for all Standards				
d.	Raw (QC)Data				
	DFTPP	NA	NA		
	Blank Data	NA	<u>NA</u>		, — /
	MS/MSD Data (if requested by USEPA Region)	NA	NA		
e.	Raw GPC Data	NA	NA		

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C/	ASE NO: 43392	SDG NO: A4B24	SDG NOs TO FOLLO	OW: N/A		
	N/A	N/A	MOD. REF. NO:	N/A		
. L	Semivolatile SIM Data	1	NA	NA		
	III-SV-SIM1 and SV-S Form VI SV-SIM; For	n II SV-SIM1 and SV-SIM2; Fo SIM2 (if required; Form IV SV-S m VII SV-SIM; Form VIII SV-SI data for QC, Samples, and St	orm SIM; IM1 and	TV V		
<u>Pe</u>	esticides Data					
a.	QC Summary					
	Surrogate Recovery Sun	nmary (Form II PEST-1 and PE	EST-2) NA	NA		
	Matrix Spike/Matrix Spike (Form III PEST-1 and PE	e Duplicate Recovery Summar EST-2)	ry <u>NA</u>	<u>NA</u>		Manager () - Close and (
	Laboratory Control Samp PEST-4)	ole Recovery (Form III PEST-3	3 and <u>NA</u>	<u>NA</u>		**Company of the Company of the Comp
	Method Blank Summary	(Form IV PEST)	NA	NA		
b. \$	Sample Data		NA	<u>NA</u>		
	TCL Results - Organics	Analysis Data Sheet (Form I P	EST)			
	Chromatograms (Primar	y Column)				
	Chromatograms from se	cond GC column confirmation				
	GC Integration report or	data system printout				
	Manual work sheets					and the same of th
	For Pesticides by GC/MS	}				
		a and copies of background-su pounds (samples & standards)				
с. 🤅	Standards Data		NA	NA		
	Initial Calibration of Sing	le Component Analytes (Form	VI PEST-1 and PEST-2)		•	-
	Toxaphene Initial Calibra	ition (Form VI PEST-3 and PE	ST-4)			
	Analyte Resolution Sum	mary (Form VI PEST-5, per co	olumn)			
	Performance Evaluation	Mixture (Form VI PEST-6)				
	Individual Standard Mixt	ure A (Form VI PEST-7)		* .		
	Individual Standard Mixt	ure B (Form VI PEST-8)				
	Individual Standard Mixt	ure C (Form VI PEST-9 and P	EST-10)			
	Calibration Verification S	ummary (Form VII PEST-1)				
	Calibration Varification S	Summary (Form VII PEST-2)				7

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HRS Reference #81 Page 60 of 126

	SE NO: 43392	SDG NO: A4B24	SDG NOs TO FO		•		
N/	'A	N/A	MOD. REF. NO	: <u> </u>	N/A		
	Calibration Verification Su	ummary (Form VII PEST-3)				M
	Calibration Verification Su	ummary (Form VII PEST-4)				
	Analytical Sequence (For	m VIII PEST)					
	Florisil Cartridge Check (F	Form IX PEST-1)				·	
	Pesticide GPC Calibration	n (Form IX PEST-2)		`			
	Identification Summary fo	or Single Component Analy	rtes (Form X PEST-1)				
	Identification Summary fo	or Toxaphene Form X PES	T-2)				
	Chromatograms and data	system printouts					
	A printout of Retentio	on Times and correspondir	ng peak areas or peak heig	ghts	•		
d.	Raw QC Data			•			
	Blank Data		<u>N</u>	IA	NA		
	Matrix Spike/Matrix Sp	pike Duplicate Data	. <u>N</u>	IA	NA		· <u> </u>
	Laboratory Control Sar	mple	<u>N</u>	IA	NA		
e.	Raw GPC Data		<u>N</u>	IA	NA		
f.	Raw Florisil Data		. <u>N</u>	IA	NA		<u> </u>
Arc	oclor Data						7
a.	QC Summary						, b
	Surrogate Recovery Si	ummary (Form II ARO-1 a	nd ARO-2) <u>1</u>	2	13		· <u> </u>
	Matrix Spike/Matrix Sp and ARO-2)	oike Duplicate Summary (F		4	17		
	Laboratory Control Sar ARO-4)	mple Recovery (Form III A		8	19		<u> </u>
	Method Blank Summa	ry (Form IV ARO)	<u>2</u>	0	21		
b.	Sample Data		· <u>2</u>	2	146		V
	TCL Results - Organic	es Analysis Data Sheet (Fo	orm I ARO) <u>N</u>	IA	NA		
	Chromatograms (Prima	ary Column)	<u></u>	IA	NA		
	Chromatograms from s	second GC column confirm	nation <u>N</u>	IA	NA		
	GC Integration report of	of data system printout	<u> </u>	IA	NA		
	Manual work sheets		<u> </u>	IA	NA		W
	For Aroclors by GC/MS	S	, N	1A	NA		

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HRS Reference #81 Page 61 of 126

CASE NO: 43392 SDG NO: A4B24	SDG NOs TO FOLL	OW: N/A		
N/A N/A	MOD. REF. NO :	N/A		·
Copies of raw spectra and copies of background-subtracted mass spectra of target compounds (samples & standards)				
Standards Data	147	318	· · · · · · · · · · · · · · · · · · ·	
Aroclors Initial Calibration (Form VI ARO-1,ARO-2, and ARO	O-3)			
Calibration Verification Summary (Form VII ARO-1)				
Analytical Sequence (Form VIII ARO)				
Identification Summary for Multicomponent Analytes (Form	X ARO)			
Chromatograms and data system printouts			7	
A printout of Retention Times and corresponding peak or peak heights	k areas			
Raw QC Data				
Blank Data	319	368		
Matrix Spike/Matrix Spike Duplicate Data	369	390		*****
Laboratory Control Sample (LCS) Data	391	402		
Raw GPC Data (if performed)	NA	NA NA		
<u>Miscellaneous Data</u>				
Original preparation and analysis forms or copies of preparat and analysis logbook pages	ion <u>403</u>	480		
Internal sample and sample extract transfer chain-of-custody records	494	495		
Screening records	NA	<u>NA</u>		
All instrument output, including strip charts from screening activities (describe or list)				
				, <u></u>
EPA Shipping/Receiving Documents				
Airbills (No. of shipments4)	481	484		
Chain of Custody Records	485	486	>	
Sample Tags				
Sample Log-in Sheet (Lab & DC-1)	49 487	7 503 493		
Miscellaneous Shipping/Receiving Records (describe or list)	· •		**************************************	

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HRS Reference #81

CASE NO: 43392	SDG NO: A4B24	SDG NOs TO F	OLLOW:	N/A		
N/A	N/A	MOD. REF. NO	: <u>N</u> /	Α		
11. Internal Lab Sample	Transfer Records and Tracking S	heets (describe or list)				
Sample Trans	sfer		494	495		
12. Other Records (des	cribe or list)					
Telephone Com	munication Log		NA	NA		
PE Ins	tructions		496	497		
						-
13. Comments						
Completed by:	Idus V Reyes	Mode			/Dco_	
(CLP Lab)	(Signature) U			e/Title)		(Date)
Verified by: (CLP Lab)	(Signature)			Preyou e/Title)	Dauti C	(Date)
Audited by:	my 2	Bi\1	lahany	Prop. Proj	scientist_	5/30/13
(USEPA)	(Signature)	(Prin	ited Nam	e/Title)		(Date)

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EPA NEW ENGLAND COMPLETE SDG FILE RECEIPT / TRANSFER FORM

Site: Jard Company Inc TOD: 12-10-0008 TASIK: 0850

		•		
Case: 43392	SDG: A44	B24		
The state of the s		en e	parameter records, and records against the second s	 and the second s

Receipt Date	Received By : Name	Init.	Affiliation	CSF Activity		dy Seals t / Intact	Released To	Date
05/09/13	Doris Guzman	25	ESAT	Received for Transfer	(Y)N	YN	Weston	05/09 / 13
5/9/13	B. Mahany	FAM	Weston	Sterge + Validation	O N	\mathcal{O}_{N}		
·	,				YN	YN		
			,		YN	YN		
					YN	YN		
					YN	YN		
					YN	YN		
					YN	YN	3.3.36	
·					YN	YN		
					YN	YN		
Page					YN	YN		
Page 64 of 12					YN	YN		•

EPA-NE - DQO SUMMARY FORM

A separate Form should be completed for each sampling event. Refer to Attachment A for instructions on completing this form, Attachment B for a complete list of the parameter codes and Attachment C for an example of a completed form.

2.	EPA Program: TSCA CERCLA RCRA Other: Projected Date(s) of Sampling Sprii EPA Site Manager Martha Bosworth EPA Case Team Members QAPP Title and Revision Date Site As Water, and Sediment Sampling Jard Approved by: Martha Bosworth Title of Approving Official: Site As	ng (April/May) h sessment Progr	ram Site Speci Bennington, V	CERCLA Sit Phase: ERA (circle one) ific Quality Ass Vermont dated of Approval:	Bennington Latitude/Le E/Spill Ident SA/SI pre- Other: Site R urance Proje I January 2 IBD	n. Vermont ongitude 42° 5: ifier NoVTD04 RI RI (phase I, eassessment ct Plan for Surf	8141741 (Inc. etc.) FS RI	lude Operable D RA post-R. urface Soil/So	Unit) A
	*If other than EPA, record date appr EPA Oversight Project (circle one) Confirmatory Analysis for Field Scre Are comparability criteria document	Y eening(Y)		Гуре of EPA O	versight (circ	le one) PRP or natory: % splits	FF Other:		
3. a.	Matrix Code ¹	SO	SO	SO	GW	GW	SD	SD	SD
ъ.	Parameter Code ²	PCB Aroclors	PCB Aroclors	PCB Congeners	PCB Aroclors	PCB Congeners	PCB Aroclors	PCB Aroclors	PCB Congeners
c.	Preservation Code ³	5	5	5	5	5	5	5	5
d.	Analytical Services Mechanism	DAS or CLP	DAS or CLP	CLP	DAS or CLP	DAS or CLP	DAS or CLP	DAS or CLP	CLP
e.	No. of Sample Locations	65	28	2	21	2	60	60	60
	Field QC:	At .							
£.	Field Duplicate Pairs	4	2		2	5	5	5	5
g.	Equipment Blanks	See RB	See RB	See RB	Şee RB	See RB	See RB	See RB	See RB
h.	VOA Trip Blanks	0	0	0	0 -	0	0	0	0
i.	Cooler Temperature Blanks	1 per cooler	l per cooler	l per cooler	1 per cooler	l per cooler	l per cooler	1 per cooler	l per cooler
j.	Bottle Blanks	0	. 0	0	0	0	0	. 0	0
k.	Other:								
l.	PES sent to Laboratory	NA	6	TBD	3	TBD	NA	3	ŤBD
	Laboratory QC:						-1	V. 1	
m.	Reagent Blank	0	0	0	0	0	0	0	0
n.	Duplicate	. 0	0	0	0	0	0	0	. 0
0.	Matrix Spike	0	2	0	1 -	0	1	0	0
p.	Matrix Spike Duplicate	0	2	0	1	0	1	0	0
q.	Other:								
4.	Site Information Site Dimensions Approximately 11. List all potentially contaminated mat Range of Dopth to Groundwater Soil Types: Surface Subsurface C Sediment Types: Stream Pond Es	trices <u>Surface</u> eater than 5 fe other:	<u>et</u> .	e soil, sediment	· · · · · · · · · · · · · · · · · · ·	er, and resident			Low

1.	EPA Program: TSCA CERCLA RCRA Other: Projected Date(s) of Sampling <u>Sprin</u> EPA Site Manager <u>Martha Bosworth</u> EPA Case Team Members	g (April/May)		Site Name Jar Site Location_ Assigned Site CERCLA Site Phase: ERA (circle one) Ot	Bennington, Latitude/Lon /Spill Identif SA/SI pre-R	Vermont gitude 42° 5 ier NoVTD04 I RI (phase	18141741 (Inc	lude Operable	Unit)
2. QAPP Title and Revision Date Site Assessment Program Site Specific Quality Assurance Project Plan for Surface and Subsurface Soil/Source, G Water, and Sediment Sampling Jard Company Inc. Bennington, Vermont dated 11 January 2013 Approved by: Martha Bosworth Date of Approval: TBD Title of Approving Official: Site Assessment Manager Organization*: EPA *If other than EPA, record date approval authority was delegated: EPA Oversight Project (circle one) Y Type of EPA Oversight (circle one) PRP or FF Other:									
	Confirmatory Analysis for Field Scree Are comparability criteria documente	ening(Y) d? Y	N II	EPA Oversight	or Confirma	tory: % splits	an n		
3. a.	Matrix Code ¹	SS	SS	SS	RB				
b.	Parameter Code ²	PCB Aroclors	PCB Aroclors	PCB Congeners	PCB Aroclors				
C.	Preservation Code ³	5	5.	5	5 .				
d.	Analytical Services Mechanism	DAS or CLP	DAS or CLP	CLP	CLP Non- RAS			N. C.	
e.	No. of Sample Locations	125	38	2	21				
	Field QC:								
f.	Field Duplicate Pairs	7	2		0				
g.	Equipment Blanks	See RB	See RB	See RB	0	-			
h.	VOA Trip Blanks	0	0 -	0	0				
i.	Cooler Temperature Blanks	1 per cooler	l per cooler	l per cooler	1 per cooler		·		
j.	Bottle Blanks	0	-0	0.	0		4		
k.	Other:							4.	
1.	PES sent to Laboratory	NA ·	6	TBD	0				
	Laboratory QC:								
m.	Reagent Blank	* 0	0	0	0				
n.	Duplicate	0	0	0	0 ·			area y properti	
0.	Matrix Spike	0	2	0	0				
p.	Matrix Spike Duplicate	0	2	0					
q.	Other:								
4.	Site Information Site Dimensions Approximately 11.1 List all potentially contaminated mat Range of Dooth to Grundwater Properties Soil Types: Surface Subsurface O Sediment Types: Stream Pond Est	rices <u>Surface</u> eater than 5 fee her:	et	e soil, sediment,				ontent: Fligh	Low

When mi	ultiple matrices will be sampled during a sam	pling event, complete Sections 5-10 for	each matrix.	Matrix Code ¹ _S	<u>SO</u>
5. Data	Engineering D	Assessment PRP Determination tent of Contamination Human and/or I esign Remedial Actio Action (quarterly monitoring)	n	Othe	ternatives
			Draft DQO Summa	ry Form 11/96	
6.	property. A subset of samples will be su	absurface soil/source samples from the ic B Aroclors field screening and fixed base abmitted for fixed laboratory analysis wit	ed laboratory analysis h a smaller subset sub	in source areas on the J mitted for PCB Conger	ard Company Inc
	Complete Table if applicable				
		·			
	COCs	Action Levels		Analytical Method-(Quantitation Limits
PCB Arc	oclors (Field Screening)	Above Background (Assumed to be N	D) 0.2	mg/Kg	<u> </u>
PCB Arc	oclors (Fixed Lab)	Above Background (Assumed to be N	D) 33 t	ıg/kg	
PCB Co	ngerners	Above Background (Assumed to be N	D) 20 t	o 100 ng/Kg	
				· · · · · · · · · · · · · · · · · · ·	
7.	Sampling Method (circle technique)	Split Spoon Dredge Trov	et or Spigot	Other:	0
	Sampling Procedures (SOP name, No., List Background Sample Locations <u>NA</u> Circle: Grab or Composite "Hot spots" sampled:	Rev. #, and date) A for source samples No			
8.	Field Data (circle) ORP pH Other:	Specific Conductance Diss		iperaturé Turb	oidity
9.	Analytical Methods and Parameters				
	Method title/SOP name	Method/SOP Identification number	Revision Date	Target Param (VOA, SV, P	eters est/PCB, Metals, etc.)
	PCB Aroclors (Field Screening)	EIA-FLDPCB2.SOP		PCBs	
	PCB Aroclors	SOM01.2 or DAS Equivalent		PCBs	
	PCB Congeners	CBC01.0		PCB Congen	ers
10.	2. Other Validation Tier (circle one)	n I, EPA-NE Data Validation Functiona Approved Validation Criteria: II Partial Tier III: ta Validation Weston Solutions, Inc./ST		ating Environmental A	
11.	Company Name <u>Weston Solutions, Inc.</u> Contract Name (e.g. START, RACS, et Person Completing Form/Title <u>G. Hon</u>		umber_EP-W-05-042 gnment No. 20114-08 Summary Form Com	1-998-0850	13

	Engineering 1	xtent of Contamination Human and/or E		Other:
		er samples from ground water monitoring waset of samples will be submitted for PCB (ells previously ins	called on and off the property for PCB Aroclors
	Complete Table if applicable			
	COCs	Action Levels		Analytical Method-Quantitation Limits
СВ А1	roclors (Fixed Lab)	Above Background (Assumed to be N	D) 1	0 μg/L
CB Co	ongeners	Above Background (Assumed to be N	D). 1	00 to 1,000 pg/L
	· · · · · · · · · · · · · · · · · · ·			
	Sampling Method (circle technique)	Bailer Low flow pump (Region I r	nethod: Yes) N	o) Peristaltic Pump
•	Sampling Procedures (SOP name, No List Background Sample Locations Circle Grab or Composite "Hot spots" sampled:	Positive Displacement Pump Split Spoon Dredge Trow	et or Spigot	Other:
	Sampling Procedures (SOP name, No List Background Sample Locations Circle Grab or Composite	Positive Displacement Pump Fauc Fauc Split Spoon Dredge Trow, Rev. #, and date) Ground Water monitoring wells TBD No	et or Spigot rel C	
	Sampling Procedures (SOP name, No List Background Sample Locations Circle Grab or Composite "Hot spots" sampled:	Positive Displacement Pump Fauc Split Spoon Dredge Trown., Rev. #, and date) Fround Water monitoring wells TBD No	et or Spigot rel C	ther:
	Sampling Procedures (SOP name, No List Background Sample Locations Circle: Grab or Composite "Hot spots" sampled: Field Data (circle) ORP pH	Positive Displacement Pump Fauc Split Spoon Dredge Trown., Rev. #, and date) Fround Water monitoring wells TBD No	et or Spigot rel C	ther:
•	Sampling Procedures (SOP name, No List Backeround Sample Locations Circle: Grab or Composite "Hot spots" sampled: Field Data (circle) ORP pH Other:	Positive Displacement Pump Fauc Split Spoon Dredge Trown., Rev. #, and date) Fround Water monitoring wells TBD No	et or Spigot rel C	emperature Turhidity
	Sampling Procedures (SOP name, No List Background Sample Locations C Circle Grab or Composite "Hot spots" sampled: Field Data (circle) ORP pH Other: Analytical Methods and Parameters	Positive Displacement Pump Split Spoon Dredge Trow "Rev. #, and date) round Water monitoring wells TBD No Specific Conductance Sisso Method/SOP	et or Spigot Co	emperature Turbidity
	Sampling Procedures (SOP name, No List Background Sample Locations C Circle Grab or Composite "Hot spots" sampled: Field Data (circle) ORP pH Other: Analytical Methods and Parameters Method title/SOP name	Positive Displacement Pump Split Spoon Dredge Trown., Rev. #, and date) Fround Water monitoring wells TBD No Specific Conductance Dissert Method/SOP Identification number	et or Spigot Co	emperature Turhidity ate Target Parameters (VOA, SV, Pest/PCB, Metals, etc.
	Sampling Procedures (SOP name, No List Background Sample Locations C Circle (Inal) or Composite "Hot spots" sampled: Field Data (circle) ORP Other: Analytical Methods and Parameters Method title/SOP name PCB Aroclors	Positive Displacement Pump Split Spoon Dredge Trow "Rev. #, and date) round Water monitoring wells TBD No Specific Conductance Dissert Method/SOP Identification number SOM01.2 or DAS Equivalent	et or Spigot Co	emperature Turbidity ate Target Parameters (VOA, SV, Pest/PCB, Metals, etc. PCBs
	Sampling Procedures (SOP name, No List Background Sample Locations C Circle Grab or Composite "Hot spots" sampled: Field Data (circle) ORP pH Other: Analytical Methods and Parameters Method title/SOP name PCB Aroclors PCB Congeners Validation Criteria (circle one) 1. Reg 2. Oth Validation Tier (circle one) 1	Positive Displacement Pump Split Spoon Dredge Trow "Rev. #, and date) round Water monitoring wells TBD No Specific Conductance Sisse Method/SOP Identification number SOM01.2 or DAS Equivalent CBC01.0	et or Spigot rel C Dived O T Revision D Guidelines for Ev.	emperature Turbidity ate Target Parameters (VOA, SV, Pest/PCB, Metals, etc. PCBs

When m	nultiple matrices will be sampled during a san	npling event, complete Sections 5-10 for e	each matrix.		Matrix Code ¹ <u>SD</u>	
5. Data	Engineering D	tent of Contamination Human and/or E	n .	ssessment	Other:	
			Draft DQO Sur	nmary Forn	a 11/96	
6. Summarize DQOs: Collect sediment samples from a wetland located west of Park Street for PCB Aroclors field screening and fixed based lat analysis. A subset of samples will be submitted for fixed laboratory analysis with a smaller subset submitted for PCB Congener analysis.						
	Complete Table if applicable					
			· · · · · · · · · · · · · · · · · · ·			
	COCs	Action Levels		Anal	ytical Method-Quantitation Limits	
PCB Ar	oclors (Field Screening)	Above Background (Assumed to be N	D)	0.2 mg/Kg		
PCB Ar	oclors (Fixed Lab)	Above Background (Assumed to be N	D)	33 ug/kg		
PCB Co	ongerners	Above Background (Assumed to be N	D) .	20 to 100 n	g/Kg	
					•	
7.	Sampling Method (circle technique)	Bailer Low flow pump (Region I r Positive Displacement Pump Fauc Split Spoon Dredge Trow	et or Spigot	<u> </u>	Peristaltic Pump Other: rect sampling	
	Sampling Procedures (SOP name, No., List Background Sample Locations W. Circle Grab or Composite "Hot spots" sampled:	Rev. #, and date)etaInd area northeast of the Jard Company	y Inc property			
3.	Field Data (circle) ORP pH	Specific Conductance Diss	olved O ₂	Temperatur	re Turbidity	
	Other:					
€.	Analytical Methods and Parameters					
	Method title/SOP name	Method/SOP Identification number	Revision l	Date	Target Parameters (VOA, SV, Pest/PCB, Metals, etc.)	
	PCB Aroclors (Field Screening)	SOM01.2			PCBs	
	PCB Aroclors	SOM01.2 or DAS Equivalent			PCBs	
	Total Metals (including Hg)	CBC01.0			PCB Congeners	
10.		on I, EPA-NE Data Validation Functiona r Approved Validation Criteria:	l Guidelines for E	valuating E	nvironmental Analyses, Part II, III or I	
	Validation Tier (circle one) I	II III Partial Tier III: ta Validation Weston Solutions, Inc./ST	ART III	Prime	or Subcontractor (circle one)	
11.	Company Name <u>Weston Solutions, Inc.</u> Contract Name (e.g. START, RACS, e	tc.) START III Work Assig	ımber_EP-W-05-	4-081-998-0	0850	
	Person Completing Form/Title_G. Hor	nok/Lead Project Scientist_Date of DQO	Summary Form (Completion	11 January 2013	

HRS Reference #81 Page 69 of 126

When multiple matrices will be sampled during a sam	pling event, complete Sections 5-10 for e	ach matrix.	. 1	Matrix Code ¹ _SS
Engineering De	ent of Contamination Human and/or E			Removal Actions Remediation Alternatives Other:
		Draft DQO Su	mmary Form	11/96
	mples from residential properties downgning and fixed based laboratory analysis in a smaller subset	n source areas o	n the Jard Co	mpany Inc property. A subset of
Complete Table if applicable				
COCs	Action Levels		Analy	tical Method-Quantitation Limits
PCB Aroclors (Field Screening)	Above Background (Assumed to be NI	D)	0.2 mg/Kg	
PCB Aroclors (Fixed Lab)	Above Background (Assumed to be NI		33 ug/kg	
PCB Congerners	Above Background (Assumed to be N		20 to 100 ng	g/Kg
				·
7. Sampling Method (circle technique) Sampling Procedures (SOP name, No., List Background Sample Locations Rec Circle: Grab or Composite "Hot spots" sampled:	Split Spoon Dredge Trow	et or Spigot el	Other: Dir	Peristaltic Pump Other:
8. Field Data (circle) ORP pH Other:	Specific Conductance Disse	olved O ₂	Temperatur	e Turbidity
Analytical Methods and Parameters				
Method title/SOP name	Method/SOP Identification number	Revision	n Date	Target Parameters (VOA, SV, Pest/PCB, Metals, etc.)
PCB Aroclors (Field Screening)	SOM01.2			PCBs.
PCB Aroclors	SOM01.2 or DAS Equivalent			PCBs
Total Metals (including Hg)	CBC01.0	المأفويل		PCB Congeners
2. Other Validation Tier (circle one) I (Company/Organization Performing Da 11. Company Name Weston Solutions, Inc. Contract Name (e.g. START, RACS, et		ART III Imber_EP-W-0 nment No. 201	Frime 5-042 14-081-998-0	or Subcontractor (circle one)
Matrix Codes ¹ - Refer to Attachment B, Part I Parameter Codes ² - Refer to Attachment B, Part II Preservation Codes ³				

K₂Cr₂O₇ Freeze Room Temperature (avoid excessive heat) Other (Specify) Not preserved



Attachment C

Original Analytical Results (Form I's) Case No. 43392; SDG No. A4B24

HRS Reference #81 Page 71 of 126

1H - FORM I ARO AROCLOR ORGANICS ANALYSIS DATA SHEET

EPA	SAMPLE	NO.	
	A4B17		-

Lab Name: Chemtech Contra	act: EPW11030	
Lab Code: CHEM Case No.: 43392 Mod. Ref	No.: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID:	E1902-13
Sample wt/vol: 30.0 (g/mL) g	Lab File ID:	PB004942.D
% Moisture: 16 Decanted:(Y/N) N	Date Received:	04/19/2013
Extraction:(Type) SOXH	Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor:	100.0
GPC Cleanup:(Y/N) N pH: 6.12	Sulfur Cleanup: (Y	//N) <u>N</u>
Acid Cleanup: (Y/N) Y		

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	3900	U
11104-28-2	Aroclor-1221	3900	U
11141-16-5	Aroclor-1232	3900	U
53469-21-9	Aroclor-1242	190000	E
12672-29-6	Aroclor-1248	3900	υ
11097-69-1	Aroclor-1254	3900	U
11096-82-5	Aroclor-1260	3900	U
37324-23-5	Aroclor-1262	3900	U
11100-14-4	Aroclor-1268	3900	U

EPA SAMPLE NO.

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.:	sDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-13DL
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004943.D
% Moisture:16 Decanted:(Y/N)_N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uI	1)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	1000.0
GPC Cleanup: (Y/N) N pH: 6.12		Sulfur Cleanup: (Y/N) <u>N</u>

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	39000	υ
11104-28-2	Aroclor-1221	39000	υ
11141-16-5	Aroclor-1232	39000	U
53469-21-9	Aroclor-1242	280000	D
12672-29-6	Aroclor-1248	39000	U
11097-69-1	Aroclor-1254	39000	U
11096-82-5	Aroclor-1260	39000	υ
37324-23-5	Aroclor-1262	39000	U
11100-14-4	Aroclor-1268	39000	U

EPA	SAMPLE	NO.

A4B18	

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.:	SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-14
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004944.D
% Moisture:6.5 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	100.0
GPC Cleanup:(Y/N) N pH: 6.05		Sulfur Cleanup: (Y/N) <u>N</u>
Agid Cleanums (V/N) V			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	3500	U
11104-28-2	Aroclor-1221	3500	U
11141-16-5	Aroclor-1232	3500	U
53469-21-9	Aroclor-1242	550000	E
12672-29-6	Aroclor-1248	3500	U
11097-69-1	Aroclor-1254	3500	υ
11096-82-5	Aroclor-1260	3500	υ
37324-23-5	Aroclor-1262	3500	υ
11100-14-4	Aroclor-1268	3500	υ

EPA SAMPLE NO.

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-14DL
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB005131.D
% Moisture:6.5 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	05/02/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	20000.0
GPC Cleanup:(Y/N) N pH: 6.05		Sulfur Cleanup: (Y/N) <u>N</u>

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	700000	υ
11104-28-2	Aroclor-1221	700000	υ
11141-16-5	Aroclor-1232	700000	U
53469-21-9	Aroclor-1242	4800000	D
12672-29-6	Aroclor-1248	700000	U
11097-69-1	Aroclor-1254	700000	U
11096-82-5	Aroclor-1260	700000	υ
37324-23-5	Aroclor-1262	700000	U
11100-14-4	Aroclor-1268	700000	U

EPA	SAMPLE	NO.
	A4B19	

Lab Name: Chemtech	Contract:	EPW11030	_
Lab Code: CHEM Case No.: 43392 Mo	d. Ref No.:	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-15
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004946.D
% Moisture: 9.8 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 10.16		Sulfur Cleanup:()	//N) <u>N</u>
Acid Cleanup: (Y/N) Y			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	37	U
11104-28-2	Aroclor-1221	37	U
11141-16-5	Aroclor-1232	37	U
53469-21-9	Aroclor-1242	770	
12672-29-6	Aroclor-1248	37	U
11097-69-1	Aroclor-1254	37	U
11096-82-5	Aroclor-1260	37	U
37324-23-5	Aroclor-1262	37	U
11100-14-4	Aroclor-1268	37	U

EPA	SAMPLE	NO.

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	_ Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-15DL
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004947.D
% Moisture:9.8 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (ul	L)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor	: 1.0	Dilution Factor:	2.0
GPC Cleanup:(Y/N) N pH: 10.16		Sulfur Cleanup: (Y/N) <u>N</u>

		CONCENTRATION UNITS:		
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q	
12674-11-2	Aroclor-1016	73	υ	
11104-28-2	Aroclor-1221	73	U	
11141-16-5	Aroclor-1232	73	U	
53469-21-9	Aroclor-1242	820	D	
12672-29-6	Aroclor-1248	73	U	
11097-69-1	Aroclor-1254	73	υ	
11096-82-5	Aroclor-1260	73	υ	
37324-23-5	Aroclor-1262	73	υ	
11100-14-4	Aroclor-1268	73	U	

EPA	SAMPLE	NO.
	A4B20	

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-16
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004948.D
% Moisture:9.3 Decanted:(Y/N) _ N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 ((uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Facto	or: 1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 8.23		Sulfur Cleanup: (Y/N) <u>N</u>

Acid	Cleanup:	(Y/N)	Y
ACIG	creamap.	(+ / + 1	-

		CONCENTRATION UNITS:		
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q	
12674-11-2	Aroclor-1016	36	U	
11104-28-2	Aroclor-1221	36	U	
11141-16-5	Aroclor-1232	36	U	
53469-21-9	Aroclor-1242	1600	E	
12672-29-6	Aroclor-1248	36	U	
11097-69-1	Aroclor-1254	36	U	
11096-82-5	Aroclor-1260	36	U	
37324-23-5	Aroclor-1262	36	U	
11100-14-4	Aroclor-1268	36	U	

EPA SAMPLE NO.
A4B20DL

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.:	sDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-16DL
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004949.D
% Moisture:9.3 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	10.0

GPC Cleanup:(Y/N) N pH: 8.23 Sulfur Cleanup:(Y/N) N

		CONCENTRATION UNITS:		
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q	
12674-11-2	Aroclor-1016	360	U	
11104-28-2	Aroclor-1221	360	U	
11141-16-5	Aroclor-1232	360	υ	
53469-21-9	Aroclor-1242	1900	D	
12672-29-6	Aroclor-1248	360	υ	
11097-69-1	Aroclor-1254	360	υ	
11096-82-5	Aroclor-1260	360	υ	
37324-23-5	Aroclor-1262	360	υ	
11100-14-4	Aroclor-1268	360	U	

EPA	SAMPLE	NO.	
	3.4D21		•

3600

3600

U

U

Lab Name: Chemtech Contract:	: EPW11030
Lab Code: CHEM Case No.: 43392 Mod. Ref No.	SDG No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID: E1902-17
Sample wt/vol: 30.1 (g/mL) g	Lab File ID: PB004950.D
% Moisture:9.0Decanted:(Y/N)N	Date Received: 04/19/2013
Extraction:(Type) SOXH	Date Extracted: 04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed: 04/26/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor: 100.0
GPC Cleanup:(Y/N) N pH: 6.48	Sulfur Cleanup:(Y/N) N

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	3600	υ
11104-28-2	Aroclor-1221	3600	υ
11141-16-5	Aroclor-1232	3600	υ
53469-21-9	Aroclor-1242	660000	E
12672-29-6	Aroclor-1248	3600	υ
11097-69-1	Aroclor-1254	3600	υ
11096-82-5	Aroclor-1260	3600	υ

Acid Cleanup: (Y/N) Y

37324-23-5

11100-14-4

Aroclor-1262

Aroclor-1268

EPA	SAMPLE	NO.

72000

U

A4B21DL

Lab Name: Chemtech	Contract:	EPW11030		
Lab Code: CHEM Case No.: 43392 Mod	d. Ref No.:	SDG	No.:	A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-	17DL
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB0051	20.D
% Moisture: 9.0 Decanted:(Y/N) N		Date Received:	04/19	/2013
Extraction:(Type) SOXH		Date Extracted:	04/19	/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	05/01	/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	200	0.0
GPC Cleanup:(Y/N) N pH: 6.48		Sulfur Cleanup:()	7/N)	N
Acid Cleanup: (Y/N) Y				

CAS NO.	COMPOUND	CONCENTRATION UNITS: (ug/L or ug/Kg) ug/Kg	Q
12674-11-2	Aroclor-1016	72000	
11104-28-2	Aroclor-1221	72000	υ
11141-16-5	Aroclor-1232	72000	υ
53469-21-9	Aroclor-1242	730000	D
12672-29-6	Aroclor-1248	72000	U
11097-69-1	Aroclor-1254	72000	U
11096-82-5	Aroclor-1260	72000	U
37324-23-5	Aroclor-1262	72000	Ŭ

11100-14-4

Aroclor-1268

EPA	SAMPLE	NO.	
	A4B22		

Lab Name: Chemtech Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392 Mod. Ref No	.: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID:	E1902-18
Sample wt/vol: 30.1 (g/mL) g	Lab File ID:	PB004957.D
% Moisture: Decanted:(Y/N) N	Date Received:	04/19/2013
Extraction:(Type) SOXH	Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 6.67	Sulfur Cleanup: (Y/N) <u>N</u>

	CONCENTRATION UNITS:		
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	36	U
11104-28-2	Aroclor-1221	36	U
11141-16-5	Aroclor-1232	36	U
53469-21-9	Aroclor-1242	12000	E
12672-29-6	Aroclor-1248	36	U
11097-69-1	Aroclor-1254	36	U
11096-82-5	Aroclor-1260	36	U
37324-23-5	Aroclor-1262	36	U
11100-14-4	Aroclor-1268	36	U

EPA	SAMPLE	NO.
	A4B22DL	

100.0

Lab	Name:	Chemtech			Contract:	EPW11030		
Lab	Code:	СНЕМ	Case No.:	43392	Mod. Ref No	.:	SDG No.:	A4B24

Matrix: (SOIL/SED/WATER) SOIL Lab Sample ID: E1902-18DL Lab File ID: PB004958.D Sample wt/vol: 30.1 (g/mL) g

% Moisture: 7.6 Decanted:(Y/N) N Date Received: 04/19/2013

Extraction:(Type) SOXH Date Extracted: 04/19/2013

Concentrated Extract Volume: 10000 (uL) Date Analyzed: 04/26/2013 Injection Volume: 1.0 (uL) GPC Factor: 1.0

GPC Cleanup:(Y/N) N pH: 6.67 Sulfur Cleanup: (Y/N) N

Acid Cleanup: (Y/N) Y

	CONCENTRATION UNITS:		
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	3600	U
11104-28-2	Aroclor-1221	3600	U
11141-16-5	Aroclor-1232	3600	U
53469-21-9	Aroclor-1242	40000	D
12672-29-6	Aroclor-1248	3600	U
11097-69-1	Aroclor-1254	3600	U
11096-82-5	Aroclor-1260	3600	U
37324-23-5	Aroclor-1262	3600	U
11100-14-4	Aroclor-1268	3600	U

Dilution Factor:

EPA	SAMPLE	NO.	
	A4B23		•

Lab Name: Chemtech Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392 Mod. Ref No	.: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID:	E1902-19
Sample wt/vol: 30.0 (g/mL) g	Lab File ID:	PB004924.D
% Moisture:15 Decanted:(Y/N)N	Date Received:	04/19/2013
Extraction:(Type) SOXH	Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 8.12	Sulfur Cleanup:()	//N) <u>N</u>
Acid Cleanup: (Y/N) Y		

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	39	U
11104-28-2	Aroclor-1221	39	U
11141-16-5	Aroclor-1232	39	U
53469-21-9	Aroclor-1242	150	
12672-29-6	Aroclor-1248	39	U
11097-69-1	Aroclor-1254	39	U
11096-82-5	Aroclor-1260	39	U
37324-23-5	Aroclor-1262	39	U
11100-14-4	Aroclor-1268	39	U

EPA	SAMPLE	NO.	
	3.4D2.4		•

Lab Name: Chemtech Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392 Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID:	E1902-01
Sample wt/vol: 30.0 (g/mL) g	Lab File ID:	PB004914.D
% Moisture:20Decanted:(Y/N) _ N	Date Received:	04/18/2013
Extraction:(Type) SOXH	Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.48	Sulfur Cleanup: (Y/N) <u>N</u>

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	41	υ
11104-28-2	Aroclor-1221	41	U
11141-16-5	Aroclor-1232	41	U
53469-21-9	Aroclor-1242	41	υ
12672-29-6	Aroclor-1248	41	υ
11097-69-1	Aroclor-1254	41	υ
11096-82-5	Aroclor-1260	41	υ
37324-23-5	Aroclor-1262	41	υ
11100-14-4	Aroclor-1268	41	U

EPA	SAMPLE	NO.
	A4B25	

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-02
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004915.D
% Moisture:11 Decanted:(Y/N)N		Date Received:	04/18/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.19		Sulfur Cleanup: (7/N) <u>N</u>

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	37	U
11104-28-2	Aroclor-1221	37	U
11141-16-5	Aroclor-1232	37	U
53469-21-9	Aroclor-1242	120	
12672-29-6	Aroclor-1248	37	U
11097-69-1	Aroclor-1254	37	U
11096-82-5	Aroclor-1260	37	ŭ
37324-23-5	Aroclor-1262	37	ŭ
11100-14-4	Aroclor-1268	37	U

EPA	SAMPLE	NO.	
	A4B26		

37

37

U

U

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392 Mod	d. Ref No.:	SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-03
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004916.D
% Moisture: 12 Decanted:(Y/N) N		Date Received:	04/18/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor: 1	1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.01		Sulfur Cleanup: (//N) <u>N</u>
Acid Cleanup: (Y/N) Y			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	37	U
11104-28-2	Aroclor-1221	37	υ
11141-16-5	Aroclor-1232	37	υ
53469-21-9	Aroclor-1242	130	
12672-29-6	Aroclor-1248	37	υ
11097-69-1	Aroclor-1254	37	υ
11096-82-5	Aroclor-1260	37	υ

37324-23-5

11100-14-4

Aroclor-1262

Aroclor-1268

HRS Reference #81 Page 87 of 126 PCB

EPA	SAMPLE	NO.	
	A4B27		

Lab Name: Chemtech Contrac	t: <u>EPW11030</u>	
Lab Code: CHEM Case No.: 43392 Mod. Ref	No.: SDG No.:	A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID: E1902	-04
Sample wt/vol:30.0 (g/mL)g	Lab File ID: PB004	917.D
% Moisture:9.2 Decanted:(Y/N) N	Date Received: 04/1	8/2013
Extraction:(Type) SOXH	Date Extracted: 04/1	9/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed: 04/2	5/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor: 1.	0
GPC Cleanup:(Y/N) N pH: 7.98	Sulfur Cleanup:(Y/N)	N

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	36	U
11104-28-2	Aroclor-1221	36	U
11141-16-5	Aroclor-1232	36	U
53469-21-9	Aroclor-1242	110	P
12672-29-6	Aroclor-1248	36	U
11097-69-1	Aroclor-1254	36	U
11096-82-5	Aroclor-1260	36	U
37324-23-5	Aroclor-1262	36	U
11100-14-4	Aroclor-1268	36	U

EPA	SAMPLE	NO.
	A4B28	

Lab Name: Chemtech Contract	EPW11030	
Lab Code: CHEM Case No.: 43392 Mod. Ref N	o.: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID:	E1902-05
Sample wt/vol: 30.0 (g/mL) g	Lab File ID:	PB004918.D
% Moisture: 9.0 Decanted:(Y/N) N	Date Received:	04/18/2013
Extraction:(Type) SOXH	Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.97	Sulfur Cleanup: (Y/N) <u>N</u>
Acid Cleanup: (Y/N) Y		

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	36	U
11104-28-2	Aroclor-1221	36	U
11141-16-5	Aroclor-1232	36	υ
53469-21-9	Aroclor-1242	140	
12672-29-6	Aroclor-1248	36	υ
11097-69-1	Aroclor-1254	36	U
11096-82-5	Aroclor-1260	36	U
37324-23-5	Aroclor-1262	36	U
11100-14-4	Aroclor-1268	36	U

EPA	SAMPLE	NO.
	A4B29	

Lab Name: Chemtech	Contract:	EPW11030	_
Lab Code: CHEM Case No.: 43392	Mod. Ref No.:	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-06
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004919.D
% Moisture: 9.4 Decanted:(Y/N) N		Date Received:	04/18/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 8.01		Sulfur Cleanup:(Y	//N) <u>N</u>
Acid Cleanup: (Y/N) Y			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	36	υ
11104-28-2	Aroclor-1221	36	U
11141-16-5	Aroclor-1232	36	υ
53469-21-9	Aroclor-1242	36	U
12672-29-6	Aroclor-1248	36	υ
11097-69-1	Aroclor-1254	36	υ
11096-82-5	Aroclor-1260	36	υ
37324-23-5	Aroclor-1262	36	υ
11100-14-4	Aroclor-1268	36	υ

EPA	SAMPLE	NO.
	A4B30	

Lab Name: Chemtech Co	ontract:	EPW11030	
Lab Code: CHEM Case No.: 43392 Mod	. Ref No.:	SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-20
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004925.D
% Moisture:11 Decanted:(Y/N)N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor: 1	.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.72		Sulfur Cleanup:()	//N) <u>N</u>
Acid Cleanup: (Y/N) Y			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	37	Ū
11104-28-2	Aroclor-1221	37	Ū
11141-16-5	Aroclor-1232	37	Ū
53469-21-9	Aroclor-1242	320	
12672-29-6	Aroclor-1248	37	ŭ
11097-69-1	Aroclor-1254	37	מ
11096-82-5	Aroclor-1260	37	Ū
37324-23-5	Aroclor-1262	37	Ū
11100-14-4	Aroclor-1268	37	Ū

EPA	SAMPLE	NO.	
	3.4D21		

Lab Name: Chemtech Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392 Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID:	E1902-21
Sample wt/vol: 30.0 (g/mL) g	Lab File ID:	PB004959.D
% Moisture:15 Decanted:(Y/N)_N	Date Received:	04/19/2013
Extraction:(Type) SOXH	Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.54	Sulfur Cleanup: (//N) <u>N</u>

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	39	U
11104-28-2	Aroclor-1221	39	U
11141-16-5	Aroclor-1232	39	υ
53469-21-9	Aroclor-1242	1200	E
12672-29-6	Aroclor-1248	39	υ
11097-69-1	Aroclor-1254	39	υ
11096-82-5	Aroclor-1260	39	υ
37324-23-5	Aroclor-1262	39	υ
11100-14-4	Aroclor-1268	39	U

EPA	SAMPLE	NO.

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.:	SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-21DL
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004960.D
% Moisture:15 Decanted:(Y/N)N	_	Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000	(uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Fact	or: 1.0	Dilution Factor:	10.0
GPC Cleanup:(Y/N) N pH: 7.54		Sulfur Cleanup:(Y/N) <u>N</u>

Acid	Cleanup:	(Y/N)	Y

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	390	υ
11104-28-2	Aroclor-1221	390	U
11141-16-5	Aroclor-1232	390	U
53469-21-9	Aroclor-1242	1600	D
12672-29-6	Aroclor-1248	390	U
11097-69-1	Aroclor-1254	390	υ
11096-82-5	Aroclor-1260	390	υ
37324-23-5	Aroclor-1262	390	υ
11100-14-4	Aroclor-1268	390	U

EPA	SAMPLE	NO.	
	A4B32		

Lab Name: Chemtech	Contract:	EPW11030	_
Lab Code: CHEM Case No.: 43392	Mod. Ref No.:	SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-22
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004961.D
% Moisture: 12 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/26/2013
<pre>Injection Volume: 1.0 (uL) GPC Factor:</pre>	1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.79		Sulfur Cleanup: (//N) <u>N</u>
Acid Cleanup: (Y/N) Y			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	38	υ
11104-28-2	Aroclor-1221	38	U
11141-16-5	Aroclor-1232	38	U
53469-21-9	Aroclor-1242	970	Е
12672-29-6	Aroclor-1248	38	U
11097-69-1	Aroclor-1254	38	υ
11096-82-5	Aroclor-1260	38	U
37324-23-5	Aroclor-1262	38	υ
11100-14-4	Aroclor-1268	38	U

EPA SAMPLE NO.

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-22DL
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004962.D
% Moisture: 12 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	10.0
GPC Cleanup:(Y/N) N pH: 7.79		Sulfur Cleanup: (Y/N) <u>N</u>
Agid Cleanups (V/N) V			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	380	υ
11104-28-2	Aroclor-1221	380	U
11141-16-5	Aroclor-1232	380	υ
53469-21-9	Aroclor-1242	1200	D
12672-29-6	Aroclor-1248	380	υ
11097-69-1	Aroclor-1254	380	υ
11096-82-5	Aroclor-1260	380	υ
37324-23-5	Aroclor-1262	380	υ
11100-14-4	Aroclor-1268	380	U

EPA	SAMPLE	NO.	
	7/1022		

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	Mod. Ref No.	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-09
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004922.D
% Moisture: 14 Decanted:(Y/N) N		Date Received:	04/18/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL	1)	Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 7.35		Sulfur Cleanup:(Y/N) <u>N</u>

		CONCENTRATION UNITS:		
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q	
12674-11-2	Aroclor-1016	38	υ	
11104-28-2	Aroclor-1221	38	U	
11141-16-5	Aroclor-1232	38	υ	
53469-21-9	Aroclor-1242	180		
12672-29-6	Aroclor-1248	38	υ	
11097-69-1	Aroclor-1254	38	υ	
11096-82-5	Aroclor-1260	38	υ	
37324-23-5	Aroclor-1262	38	υ	
11100-14-4	Aroclor-1268	38	υ	

EPA	SAMPLE	NO.	
	7 / D 2 /		

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: <u>CHEM</u> Case No.: <u>43392</u> N	Mod. Ref No.:	SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-23
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004926.D
% Moisture: 14 Decanted:(Y/N) N		Date Received:	04/19/2013
Extraction:(Type) SOXH		Date Extracted:	04/22/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: 5.40		Sulfur Cleanup: (//N) <u>N</u>
Acid Cleanup: (Y/N) Y			

		CONCENTRATION UNITS:		
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q	
12674-11-2	Aroclor-1016	38	U	
11104-28-2	Aroclor-1221	38	U	
11141-16-5	Aroclor-1232	38	U	
53469-21-9	Aroclor-1242	280	P	
12672-29-6	Aroclor-1248	38	U	
11097-69-1	Aroclor-1254	38	U	
11096-82-5	Aroclor-1260	38	U	
37324-23-5	Aroclor-1262	38	U	
11100-14-4	Aroclor-1268	38	U	

EPA	SAMPLE	NO.

2	44B35
-	11000

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392	_ Mod. Ref No.:	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-10
Sample wt/vol:30.0 (g/mL)g		Lab File ID:	PB004923.D
% Moisture: <u>14</u> Decanted:(Y/N) N		Date Received:	04/18/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (un	L)	Date Analyzed:	04/25/2013
Injection Volume: 1.0 (uL) GPC Factor	: 1.0	Dilution Factor:	1.0
GPC Cleanup: (Y/N) N pH: 7.92		Sulfur Cleanup: (Y/N) <u>N</u>

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	38	U
11104-28-2	Aroclor-1221	38	U
11141-16-5	Aroclor-1232	38	υ
53469-21-9	Aroclor-1242	38	υ
12672-29-6	Aroclor-1248	38	υ
11097-69-1	Aroclor-1254	38	υ
11096-82-5	Aroclor-1260	38	υ
37324-23-5	Aroclor-1262	38	υ
11100-14-4	Aroclor-1268	38	U

EPA	SAMPLE	NO.	
	24B56		

Lab Name: Chemtech	Contract:	EPW11030		
Lab Code: CHEM Case No.: 43392	Mod. Ref No.:	SDG	No.: A4B	24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-11	
Sample wt/vol: 30.0 (g/mL) g		Lab File ID:	PB004938.D)
% Moisture:0.0 Decanted:(Y/N) N		Date Received:	04/18/201	.3
Extraction:(Type) SOXH		Date Extracted:	04/19/201	.3
Concentrated Extract Volume: _10000 (uL))	Date Analyzed:	04/25/201	.3
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	1.0	<u> </u>

GPC Cleanup:(Y/N) N pH: N/A Sulfur Cleanup:(Y/N) N

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	33	U
11104-28-2	Aroclor-1221	33	U
11141-16-5	Aroclor-1232	33	U
53469-21-9	Aroclor-1242	770	E
12672-29-6	Aroclor-1248	33	U
11097-69-1	Aroclor-1254	33	U
11096-82-5	Aroclor-1260	33	U
37324-23-5	Aroclor-1262	33	U
11100-14-4	Aroclor-1268	33	U

EPA	SAMPLE	NO.

A4B56DL

Lab Name:	Chemtech			Coı	ntract:	EPW11030		
Lab Code:	CHEM	Case No.:	43392	Mod.	Ref No.:	sDG	No.: 2	A4B24
Matrix: (S	SOIL/SED/WA	TER) SOIL				Lab Sample ID:	E1902-1	lDL
Sample wt/	/vol:30.0	(g/mL)	g			Lab File ID:	PB00493	9.D
% Moisture	e: <u>0.0</u> 1	Decanted: (Y	7/N) <u>N</u>			Date Received:	04/18/	2013
Extraction	n:(Type) SO	хн				Date Extracted:	04/19/	2013
Concentrat	ted Extract	Volume:	10000 (uI	۲)		Date Analyzed:	04/25/	2013
Injection	Volume: 1.	0 (uL)	GPC Factor	1.0	0	Dilution Factor:	5.0	
GPC Clean	up:(Y/N) N	pH: _1	1/A			Sulfur Cleanup: (Y/N)	<u>N</u>
344 0144	(37/NT)	37						

Acid	Cleanup:	(Y/N)	Y
11014	crcanap.	(+ / + 1 /	_

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	160	U
11104-28-2	Aroclor-1221	160	U
11141-16-5	Aroclor-1232	160	U
53469-21-9	Aroclor-1242	870	D
12672-29-6	Aroclor-1248	160	U
11097-69-1	Aroclor-1254	160	υ
11096-82-5	Aroclor-1260	160	υ
37324-23-5	Aroclor-1262	160	υ
11100-14-4	Aroclor-1268	160	U

EPA	SAMPLE	NO.	
	3.4DE7		

Lab Name: Chemtech Contract	t: <u>EPW11030</u>	
Lab Code: CHEM Case No.: 43392 Mod. Ref 1	No.: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL	Lab Sample ID:	E1902-12
Sample wt/vol: 30.1 (g/mL) g	Lab File ID:	PB004940.D
% Moisture:O.O Decanted:(Y/N)N	Date Received:	04/18/2013
Extraction:(Type) SOXH	Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)	Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor: 1.0	Dilution Factor:	1.0
GPC Cleanup:(Y/N) N pH: N/A	Sulfur Cleanup:(Y/N) <u>N</u>

·		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	33	U
11104-28-2	Aroclor-1221	33	U
11141-16-5	Aroclor-1232	33	U
53469-21-9	Aroclor-1242	33	U
12672-29-6	Aroclor-1248	33	U
11097-69-1	Aroclor-1254	33	U
11096-82-5	Aroclor-1260	810	Е
37324-23-5	Aroclor-1262	33	υ
11100-14-4	Aroclor-1268	33	U

EPA	SAMPLE	NO.

A4B57DI	<u>.</u>

Lab Name: Chemtech	Contract:	EPW11030	
Lab Code: CHEM Case No.: 43392 M	od. Ref No.:	: SDG	No.: A4B24
Matrix: (SOIL/SED/WATER) SOIL		Lab Sample ID:	E1902-12DL
Sample wt/vol: 30.1 (g/mL) g		Lab File ID:	PB004941.D
% Moisture:0.0 Decanted:(Y/N)N		Date Received:	04/18/2013
Extraction:(Type) SOXH		Date Extracted:	04/19/2013
Concentrated Extract Volume: 10000 (uL)		Date Analyzed:	04/26/2013
Injection Volume: 1.0 (uL) GPC Factor:	1.0	Dilution Factor:	5.0
GPC Cleanup:(Y/N) N pH: N/A		Sulfur Cleanup: (Y	//N) <u>N</u>
Acid Cleanup: (Y/N) Y			

		CONCENTRATION UNITS:	
CAS NO.	COMPOUND	(ug/L or ug/Kg)ug/Kg	Q
12674-11-2	Aroclor-1016	160	υ
11104-28-2	Aroclor-1221	160	U
11141-16-5	Aroclor-1232	160	U
53469-21-9	Aroclor-1242	160	U
12672-29-6	Aroclor-1248	160	υ
11097-69-1	Aroclor-1254	160	U
11096-82-5	Aroclor-1260	830	D
37324-23-5	Aroclor-1262	160	U
11100-14-4	Aroclor-1268	160	Ū



Attachment D

USEPA Contract Laboratory Program Statement of Work for Organic Analysis,
Multi-Media Multi-Concentration, SOMO1.2 (Excerpt)
and
Modifications Updating SOM01.1 to SOM01.2, October 5, 2006, Updated 02-12-2007,
Amended 04-11-2007 (Excerpt)

HRS Reference #81 Page 103 of 126

4.0 AROCLORS TARGET COMPOUND LIST AND CONTRACT REQUIRED QUANTITATION LIMITS1

			Quantitati	Quantitation Limits	
			Water	Soil	
Aroclors		CAS Number	ug/L	ug/kg	
141. 142. 143. 144. 145.	Aroclor-1016 Aroclor-1221 Aroclor-1232 Aroclor-1242 Aroclor-1248	12674-11-2 11104-28-2 11141-16-5 53469-21-9 12672-29-6	1.0 1.0 1.0 1.0	33 33 33 33 33	
146. 147. 148. 149.	Aroclor-1254 Aroclor-1260 Aroclor-1262 Aroclor-1268	11097-69-1 11096-82-5 37324-23-5 11100-14-4	1.0 1.0 1.0	33 33 33 33	

 $^{^{\}rm 1}{\rm There}$ is no differentiation between the preparation of low and medium soil samples in this method for the analysis of Aroclors.

	EXHIBIT D – AROCLORS
EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 1 Exhibit D - Aroclor: Section 7.2.3.4.1	The following Section:
Eximult D - Arociot. Section 7.2.3.4.1	"Prepare five-point initial calibration standard solutions containing a mixture of Aroclors 1016 and 1260 at the following suggested levels: 100; 200; 400; 800; and 1600 ng/mL and surrogates at 5.0, 10, 20, 40 and 80 ng/mL for tetrachloro-m-xylene and 10, 20, 40, 80 and 160 ng/mL for decachlorobiphenyl. Also, prepare a single-point initial calibration standard solution containing Aroclors 1221, 1232, 1242, 1248, 1254, 1262, and 1268 at 400 ng/mL and surrogates at 20 ng/mL for tetrachloro-m-xylene and 40 ng/mL for decachlorobiphenyl. The solutions must be prepared every 6 months, or sooner if the solutions have degraded or concentrated."
	Is updated to: "Prepare five-point initial calibration standard solutions containing a mixture of Aroclors 1016 and 1260 at the following suggested levels: 100; 200; 400; 800; and 1600 ng/mL and surrogates at 5.0, 10, 20, 40 and 80 ng/mL for tetrachloro-m-xylene and 10, 20, 40, 80 and 160 ng/mL for decachlorobiphenyl. In addition, prepare a single-point initial calibration standard solution containing Aroclors 1221 at 400 ng/mL including surrogates, tetrachloro-m-xylene at 20 ng/mL and decachlorobiphenyl at 40 ng/mL. Also, prepare a single point calibration initial calibration standard of Aroclor 1232, 1242, 1248, 1254, 1262, and 1268 as instructed for Aroclor 1221. Refer to Section 7.2.3.4.3 for five-point calibration standards of the other Aroclors. The solutions must be prepared every 6 months, or sooner if the solutions have degraded or concentrated."
Aro-Item 2 Exhibit D - Aroclor: Section 7.2.3.4.2	The following Section: "Prepare a single-point calibration verification standard solution containing Aroclor 1260 and Aroclor 1016 at 400 ng/mL and surrogates at 20 ng/mL for tetrachloro-m-xylene and 40 ng/mL for decachlorobiphenyl. The solution must be prepared every 6 months, or sooner if the solution has degraded or concentrated."
	Is updated to: "Prepare a single-point calibration verification standard solution containing Aroclor 1260 and Aroclor 1016 at 400 ng/mL and surrogates, tetrachloro-m-xylene at 20 ng/mL and decachlorobiphenyl 40 ng/mL. Additional individual calibration verification standard solution(s) containing any other Aroclor may be prepared when necessary at 400 ng/mL, including surrogates, tetrachloro-m-xylene at 20 ng/mL and decachlorobiphenyl at 40 ng/mL. The solution must be prepared every 6 months, or sooner if the solution has degraded or concentrated."

EXHIBIT/SECTION(S) MODIFICATION (S) The following Section: Aro-Item 3 Exhibit D - Aroclor: Section 9.2.1 "Summary of Initial Calibration Prior to sample analysis (including LCSs and MS/MSDs) and required blanks (method/sulfur cleanup/instrument), each GC/ECD system must be initially calibrated to determine instrument sensitivity and the linearity of Aroclor response. An initial five-point calibration is performed using Aroclors 1016 and 1260 to demonstrate the linearity of the detector response. The other seven Aroclors are calibrated at a single mid-point for pattern recognition. The standards for these seven Aroclors should be analyzed before the analysis of any samples, and may be analyzed before or after the analysis of the five levels of the Aroclor 1016/1260 standards. is updated to: Summary of Initial Calibration Prior to sample analysis (including LCSs and MS/MSDs) and required blanks (method/sulfur cleanup/instrument), each GC/ECD system must be initially calibrated to determine instrument sensitivity and the linearity of Aroclor response. An initial five-point calibration is performed using Aroclors 1016 and 1260 to demonstrate the linearity of the detector response. The other seven Aroclors can be calibrated at a single mid-point at a minimum, for pattern recognition. The standards for these seven Aroclors should be analyzed before the analysis of any samples, and may be analyzed before or after the analysis of the five levels of the Aroclor 1016/1260 standards. Note: All Aroclor target compounds may have five-point calibrations performed initially, prior to sample analyses. Alternately, as long as a valid five-point calibration of Aroclor 1016/1260 is present, five-point calibrations for any of the remaining Aroclor target compounds may be performed, prior to

sample analyses.

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 4	The following Section:
Aro-Item 4 Exhibit D - Aroclor: Section 9.2.2	Each GC/ECD system must be initially calibrated upon award of the contract, whenever major instrument maintenance or modification is performed (e.g., column replacement or repair, cleaning or replacement of the ECD, etc.), or if the calibration verification technical acceptance criteria have not been met. Also, for any sample in which an Aroclor, other than Aroclor 1016 or Aroclor 1260 is detected, results for the specific Aroclor(s) may only be reported if the Aroclor(s) have been calibrated using multipoint standards (five-point). If time remains in the 12-hour period after a valid five-point initial calibration for a detected Aroclor(s) has been performed, then samples containing the Aroclor(s) may be analyzed. If the previously-analyzed five-point initial calibration containing the Aroclor(s) detected in the sample(s) is not in the same 12-hour sequence, then the sample(s) must be analyzed after a Continuing Calibration Verification (CCV) analysis containing the Aroclor(s) detected in the sample(s) that meets the criteria for CCVs in Section 9.3.
	is updated to: Each GC/ECD system must be initially calibrated upon award of the contract, whenever major instrument maintenance or modification is performed (e.g., column replacement or repair, cleaning or replacement of the ECD, etc.), or if the calibration verification technical acceptance criteria have not been met. Also, for any sample, in which an Aroclor (other than Aroclor 1016 or Aroclor 1260) is detected, for which a valid five point calibration curve is not available, results for these specific Aroclors must be reported as an estimated concentration with the appropriate compound qualifier. Subsequently, the sample must be re-analyzed following a valid five point calibration of the specific Aroclor. All sample analysis, must be preceded by an opening CCV with an Aroclor 1016/1260 CS3 standard, at a minimum. Additional Aroclor opening CCV standards may be analyzed at the laboratory's discretion. The closing CCV must include Aroclor 1016/1260 CS3 and all detected Aroclors in the sample. When an Aroclor, other than Aroclor 1016/1260, is detected in a sample, the closing CCV CS3 standard of this detected Aroclor standard must meet opening CCV technical acceptance
Aro-Item 5 Exhibit D – Aroclor: Section 9.2.3.3	criteria in Section 9.3.5, if the sample was not preceded by the Aroclor included as a CS3 standard in the opening CCV." The following Section: "If Aroclors other than Aroclor 1016/1260 are detected in an analysis, a
	is updated to: "If Aroclors other than Aroclor 1016/1260 are detected in a sample analysis, following a single-point calibration must be prepared (Section 7.2.3.4.3) and run for that particular Aroclor, a separate five-point calibration must be prepared (Section 7.2.3.4.3) and run for that particular Aroclor, followed by a re-analysis of the sample."

EXHIBIT/SECTION(S)	MODIFICATION (S)	
EXHIBIT/SECTION(S)	MODIFICATION (S)	
Aro-Item 6	Analyze the initial calibration sequence as given below.	
Exhibit D – Aroclor: Section 9.2.3.5		
	Initial Calibration Sequence	
	1. Aroclor 1221 CS3 (400 ng/mL)	
	2. Aroclor 1232 CS3 (400 ng/mL)	
	3. Aroclor 1242 CS3 (400 ng/mL)	
	4. Aroclor 1248 CS3 (400 ng/mL)	
	5. Aroclor 1254 CS3 (400 ng/mL)	
a second control of the second control of th	6. Aroclor 1262 CS3 (400 ng/mL)	
·	7. Aroclor 1268 CS3 (400 ng/mL)	
	8. Aroclor 1016/1260 CS1 (100 ng/mL)	
	9. Aroclor 1016/1260 CS2 (200 ng/mL)	
	10. Aroclor 1016/1260 CS3 (400 ng/mL)	
	11. Aroclor 1016/1260 CS4 (800 ng/mL)	
	12. Aroclor 1016/1260 CS5 (1600 ng/mL)	
	13. Instrument blank	
	Note: The single-point Aroclor standards may be analyzed	
	after the analysis of the five levels of the Aroclor 1016/1260	
	standards. The steps pertaining to the instrument blank are	
•	used as part of the calibration verification as well.	
	asou as part of the canonation vortification as work.	
•		
	is updated to:	
	"Initial Calibration may be performed by any of the following	
	sequence Options given below:	
	Initial Calibration Sequence – Option 1	
	1. Aroclor 1221 CS3 (400 ng/mL)	
•	2. Aroclor 1232 CS3 (400 ng/mL)	
	3. Aroclor 1242 CS3 (400 ng/mL)	
	4. Aroclor 1248 CS3 (400 ng/mL)	
	5. Aroclor 1254 CS3 (400 ng/mL)	
	6. Aroclor 1262 CS3 (400 ng/mL)	
	7. Aroclor 1268 CS3 (400 ng/mL)	
	• • • • • • • • • • • • • • • • • • • •	
	9. Aroclor 1016/1260 CS2 (200 ng/mL)	
	10. Aroclor 1016/1260 CS3 (400 ng/mL)	
the state of the s	11. Aroclor 1016/1260 CS4 (800 ng/mL)	
	12. Aroclor 1016/1260 CS5 (1600 ng/mL)	
	Note: The single-point Aroclor standards may be analyzed	
	after the analysis of the five levels of the Aroclor 1016/1260	
	standards in Option 1 above.	
	OR	

HRS Reference #81

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 6	Initial Calibration Sequence - Option 2
Exhibit D – Aroclor: Section 9.2.3.5	5-points of Aroclor 1016/1260(100ng/mL to 1600ng/mL)
(Cont.)	5-points of Aroclor 1221 (100ng/mL to 1600ng/mL)
	5-points of Aroclor 1232(100ng/mL to 1600ng/mL)
	5-points of Aroclor 1242(100ng/mL to 1600ng/mL)
	5-points of Aroclor 1248(100ng/mL to 1600ng/mL)
	5-points of Aroclor 1254(100ng/mL to 1600ng/mL)
	5-points of Aroclor 1262(100ng/mL to 1600ng/mL)
	5-points of Aroclor 1268(100ng/mL to 1600ng/mL)
	OR
	Initial Calibration Sequence - Option 3
	5-points of Aroclor 1016/1260(100ng/mL to 1600ng/mL)
	5-points or single point Aroclor 1221 (100ng/mL - 1600ng/mL or 400ng/mL)
	5-points or single point Aroclor 1232 (100ng/mL - 1600ng/mL or 400ng/mL)
	5-points or single point Aroclor 1242 (100ng/mL - 1600ng/mL or 400ng/mL)
	5-points or single point Aroclor 1248 (100ng/mL - 1600ng/mL or 400ng/mL)
	5-points or single point Aroclor 1254 (100ng/mL - 1600ng/mL or 400ng/mL)
	5-points or single point Aroclor 1262 (100ng/mL- 1600ng/mL or 400ng/mL)
	5-points or single point Aroclor 1268 (100ng/mL - 1600ng/mL or 400ng/mL)
	Note: Option 2 and 3 Initial Calibration above may be performed in any Aroclor sequence as long as a valid five-point calibration of Aroclor 1016/1260 is present. Refer to Section 7.2.3.4 for initial calibration standard concentrations.

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 7	The following Section:
Exhibit D – Aroclor: Section 9.2.4.2	"For Aroclors 1016 and 1260, an RT is measured for a minimum of 3 peaks in each of the five calibration standards and the mean RT (RT) is calculated for each of the peaks as the average of the five values obtained from the five calibration standards. For Aroclors 1221, 1232, 1242, 1248, 1254, 1262, and 1268 an RT is measured for each of the peaks for a single-point calibration standard. If a valid five-point calibration is present for a specific Aroclor then an RT is measured for each of the peaks in each of the five calibration standards and the RT is calculated as the average of the five values for each of the peaks obtained from the five calibration standards. An RT is measured for the surrogates in each of the five calibration standards and the RT is calculated as the average of the five values. Calculate the RT using Equation 1:
•	is updated to:
	"For Aroclors 1016 and 1260, an RT is measured for a minimum of 3 peaks in each of the five calibration standards and the mean RT (RT) is
	calculated for each of the peaks as the average of the five values obtained from the five calibration standards. For Aroclors 1221, 1232,
	1242, 1248, 1254, 1262, and 1268 an RT is measured for a minimum
	of three peaks for a single-point calibration standard. If a valid five- point calibration is present for a specific Aroclor then an RT is measured for a minimum of three peaks in each of the five calibration
	standards and the RT is calculated as the average of the five values for each of the peaks obtained from the five calibration standards. An RT is measured for the surrogates in each of the five calibration standards
	of Aroclor 1016/1260, or from Aroclor 1016 if analyzed as a
	separate mixture. The surrogate RT is calculated as the average of
	the five values. Calculate the RT using Equation 1:"

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 8	The following Section:
Exhibit D – Aroclor: Section 9.2.4.4	"The linearity of the instrument is determined by calculating a Percent Relative Standard Deviation (%RSD) of the Calibration Factors (CFs). Either peak area or peak height may be used to calculate CFs used in the %RSD equation.
	Five sets of CFs will be generated for the Aroclor 1016/1260 mixture, each set consisting of the CFs for each of the five peaks chosen for this mixture. The single standard for each of the other Aroclors will generate at least three CFs, one for each selected peak, unless a valid five-point calibration is present for a specific Aroclor, in which case five sets of CFs will be generated for the specific Aroclor.
	Calculate CFs, the Mean CF (CF), and the %RSD of the CFs for each peak in a selected set of a minimum of 3 major peaks for each Aroclor using Equations 2, 3, and 4."
	Is updated to:
	"The linearity of the instrument is determined by calculating a Percent Relative Standard Deviation (%RSD) of the Calibration Factors (CFs). Either peak area or peak height may be used to calculate CFs used in the %RSD equation.
	Five sets of CFs will be generated for the Aroclor 1016/1260 mixture, each set consisting of the CFs for each of the peaks (minimum of three) chosen for this mixture. The single standard for each of the other Aroclors will generate at least three CFs, one for each selected peak, unless a valid five-point calibration is present for a specific Aroclor, in which case five sets of CFs will be generated for the specific Aroclor. Calibration Factors (CF) for the surrogates must be generated for each of the five calibration standards of Aroclor 1016/1260, or from Aroclor 1016 if analyzed as a separate mixture.
	The \overline{CF} of each surrogate compound is calculated as the average of the five values.
	Calculate CFs, the Mean CF (CF), and the %RSD of the CFs for each peak in a selected set of a minimum of 3 major peaks for each Aroclor using Equations 2, 3, and 4."

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 9	The following Section:
Exhibit D – Aroclor: Section 9.3.1	"Summary of Continuing Calibration Verification (CCV)
	The analyses of instrument blanks and the required Aroclor CS3 Standard Mixtures (see Section 9.3.2) constitute the calibration verification. Sample (including LCS and MS/MSD) and required blank (method/sulfur cleanup) data are not acceptable unless bracketed by acceptable analyses of instrument blanks and the Aroclor CS3 Standard Mixtures. In cases where a valid five-point initial calibration for the detected Aroclors is required, that initial calibration may be substituted for the opening CCV."
	Is updated to:
	"Summary of Continuing Calibration Verification (CCV)
	The analyses of instrument blanks and the required Aroclor CS3 Standard Mixtures (see Section 9.3.2) constitute the calibration verification. Sample (including LCS and MS/MSD) and required blank (method/sulfur cleanup) data are not acceptable unless bracketed by acceptable analyses of instrument blanks and the Aroclor CS3 Standard Mixtures."
	Note the last sentence in the section is deleted:
	"In cases where a valid five-point initial calibration for the detected Aroclors is required, that initial calibration may be substituted for the opening CCV."

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 10	The following section:
Exhibit D – Aroclor: Section 9.3.2.1	An instrument blank and Aroclor 1016/1260 CS3 Standard Mixture must bracket one end of a 12-hour period (opening CCV) during which sample and required blank data are collected, and a second instrument blank and the Aroclor 1016/1260 CS3 Standard Mixture must bracket the other end of the 12-hour period (closing CCV). If during any 12-hour period, an Aroclor other than 1016 or 1260 is detected and the 12-hour time period for the five-point initial calibration of the detected Aroclor(s) has elapsed, then an instrument blank and a CS3 standard of the detected Aroclor(s) must bracket both ends of the 12-hour period. If the opening CCV does not meet all technical acceptance criteria, then
	a new valid five-point initial calibration for the detected Aroclors must be performed before samples containing the detected Aroclors may be analyzed. is updated to:
	"An instrument blank and Aroclor 1016/1260 CS3 Standard Mixture must bracket one end of a 12-hour period (opening CCV) during which sample and required blank data are collected, a second instrument blank, Aroclor 1016/1260 CS3 and CS3 Standard Mixture (s) of any other detected Aroclor (s) must bracket the other end of a 12-hour period (closing CCV). Each opening CCV must include an instrument blank and Aroclor 1016/1260 CS3 standard, additional Aroclor CS3 standards may be performed at the laboratory's discretion. If a valid five-point calibration is available for Aroclor
	(s) other than 1016/1260, an opening CCV with an instrument blank and Aroclor 1016/1260 CS3 is sufficient, however, the closing CCV must <i>include</i> all Aroclors detected and meet opening CCV technical acceptance criteria in Section 9.3.5.3.

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 11	For the 12-hour period immediately following the initial calibration
Exhibit D – Aroclor: Section 9.3.2.2	sequence, the instrument blank is the last step in the initial calibration
	sequence and brackets the front end of that 12-hour period. The
	injection of the instrument blank starts the beginning of the 12-hour period (Section 10.3.2.1.1), followed by the injection of the Aroclor
	1016/1260 CS3 Standard. Samples (including LCSs and MS/MSDs)
	and required blanks (method/sulfur cleanup) may be injected for 12
	hours from the injection of the instrument blank. The first injections
· · · · · · · · · · · · · · · · · · ·	immediately after that 12-hour period must be an instrument blank and
	the Aroclor 1016/1260 CS3 Standard Mixture. The instrument blank must be analyzed first, before the standard.
	inust be analyzed inst, before the standard.
	Is updated to:
	"The injection of an instrument blank starts the beginning of the 12-
	hour period (Section 10.3.2.1.1), followed by the injection of Aroclor
·	1016/1260 CS3 Standard and any additional CS3 Standard
	Mixture(s) as determined by the laboratory. Samples (including
	LCSs and MS/MSDs) and required blanks (method/sulfur cleanup) may
	be injected for 12 hours from the injection of the instrument blank. The first injections immediately after the previous 12-hour period must be
	an instrument blank, Aroclor 1016/1260 CS3 Standard and CS3
	Standard Mixture(s) of any other detected Aroclor. A closing CCV
	must bracket the end of a 12-hour sequence.

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 12	The following Section:
Exhibit D – Aroclor: Section 9.3.2.3	"The analyses of the instrument blank and CS3 Standard Mixture (closing CCV) immediately following one 12-hour period may be used to begin the subsequent 12-hour period as an opening CCV, provided that they meet the technical acceptance criteria in Section 9.3.5. In that instance, the subsequent 12-hour period must be bracketed by the acceptable analyses of an instrument blank and a CS3 Standard Mixture (closing CCV), in that order. Those two analyses may in turn be used to bracket the front end of yet another 12-hour period (opening CCV). This progression may continue every 12 hours until such time as any of the instrument blanks or the CS3 Standard Mixture fails to meet the technical acceptance criteria in Section 9.3.4, or an Aroclor has been detected in a sample for which the corresponding CS3 standard was not performed for the opening CCV. The 12-hour time period begins with
	the injection of the instrument blank."
	is updated to:
	"The analyses of the instrument blank and CS3 Standard Mixture(s) (closing CCV) immediately following one 12-hour period may be used to begin the subsequent 12-hour period as an opening CCV, provided that they meet the technical acceptance criteria in Section 9.3.5. In that instance, the subsequent 12-hour period must be bracketed by the
	acceptable analyses of an instrument blank and a CS3 Standard Mixture(s) (closing CCV), in that order. Those two analyses may in turn be used to bracket the front end of yet another 12-hour period (opening CCV). This progression may continue every 12 hours until such time as any of the instrument blanks or the required CS3 Standard
	Mixture (s) fails to meet the technical acceptance criteria in Section 9.3.5.
Aro-Item 13	The following section is deleted:
Exhibit D – Aroclor: Section 9.3.2.4	"If more than 12 hours have elapsed since the injection of the instrument blank that bracketed a previous 12-hour period, an acceptable instrument blank and an Aroclor 1016/1260 CS3 standard must be analyzed in order to start a new sequence. This requirement applies even if no analyses were performed since that standard was injected."

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 14	The following Section:
Exhibit D – Aroclor: Section 9.3.2.5	"The requirements for running the instrument blanks and CS3 Aroclor 1016/1260 Standard Mixture are waived when no samples (including LCSs and MS/MSDs), dilutions, reanalyses, or required blanks (method/sulfur cleanup) are analyzed during that 12-hour period. To resume analysis, using the existing initial calibration, the Contractor must first analyze an instrument blank and CS3 Aroclor 1016/1260 Standard that meet the technical acceptance criteria."
	Is updated to:
	"The requirements for running the instrument blanks and CS3 Aroclor 1016/1260 Standard Mixture are waived when no samples (including LCSs and MS/MSDs), dilutions, reanalyses, or required blanks (method/sulfur cleanup) are analyzed during that 12-hour period. To resume analysis, using the existing initial calibration, the Contractor must first analyze an opening CCV that consist of an instrument blank, Aroclor 1016/1260 CS3 Standard, and any additional CS3 Aroclor Standard (s) that meet the technical acceptance criteria. Note: Additional opening CCV CS3 Aroclor Standard (s) determined to be necessary are at the laboratory's discretion."
Aro-Item 15 Exhibit D – Aroclor: Section 9.3.2.5	The current "Section 9.3.2.5" is updated to "Section 9.3.2.4".
Aro-Item 16	The following Section:
Exhibit D – Aroclor: Section 9.3.2.6	"If the entire 12-hour period is not required for the analyses of all samples and blanks to be reported and all data collection is to be stopped, the sequence must be ended with the instrument blank/CS3 Aroclor Standard Mixture (s) (1016/1260 and all detected Aroclors) combination."
	is updated to:
	"If the entire 12-hour period is not required for the analyses of all samples and blanks to be reported and all data collection is to be stopped, the sequence must end with an appropriate closing CCV combination, that is, an instrument blank/CS3 Aroclor 1016/1260 and all detected Aroclor CS3 Standard Mixture(s)."
Aro-Item 17	The current "Section 9.3.2.6" is updated to "Section 9.3.2.5".
Exhibit D – Aroclor: Section 9.3.2.6 Aro-Item 18	The following Section:
Exhibit D – Aroclor: Section 9.3.2.7	The following Section: "No more than 14 hours may elapse from the injection beginning the opening CCV (instrument blank) and the injection ending the closing CCV (Aroclor Standard)."
	Is updated to: "No more than 14 hours may elapse from the injection beginning the opening CCV (instrument blank) and the injection ending the closing CCV (Aroclor Standard). If more than 12 hours elapse between the injections of the two instrument blanks (opening and closing CCV) that bracket a 12-hour period in which samples or required blanks are analyzed, then the time between the injection of the instrument blank (closing CCV) and the preceding sample may not exceed the length of one chromatographic run."

Aro-Item 19	The current "Section 9.3.2.7" is updated to "Section 9.3.2.6".
Exhibit D – Aroclor: Section 9.3.2.7	·
Aro-Item 20	The following Section:
Exhibit D – Aroclor: Section 9.3.4	"Calculations for Calibration Verification
	For each analysis of the CS3 Individual Standard Mixture(s) used to demonstrate calibration verification, calculate the Percent Difference between the CF of each Aroclor peak (including the surrogates) in the standard mixture and the CF from the initial calibration, using Equation 5."
	is updated to: "Calculations for Calibration Verification
	Calculations for Calibration Verification
	For each analysis of the CS3 Individual Standard Mixture(s) used to demonstrate calibration verification, calculate the Percent Difference between the CF of each Aroclor peak in the standard mixture and the
	CF from the initial calibration, using Equation 5. Calculate the Percent Difference between CF of surrogates in each standard mixture and the CF from the initial calibration of Aroclor 1016/1260 or 1016 if analyzed as a separate mixture, using Equation 5."

EXHIBIT/SECTION(S)	MODIFICATION (S)		
Aro-Item 21	The following Section:		
Exhibit D — Aroclor: Section 9.3.5.3	"For the opening CCV, Percent Difference for each Aroclor peak and surrogates calculated from the CCV standard must not exceed $\pm 15\%$. For the closing CCV, Percent Difference for each Aroclor peak and surrogates calculated from the CCV must not exceed $\pm 50\%$. If the Percent Difference for the closing CCV is $\pm 15\%$ or less, then it can be used for the opening CCV of the next 12-hour period."		
	is updated to:		
	"For the opening CCV, Percent Difference for each Aroclor peak and surrogates calculated from the CCV standard must not exceed ±15%. For the closing CCV, Percent Difference for each Aroclor peak and surrogates calculated from the CCV must not exceed ±50%. If the Percent Difference for the closing CCV is ±15% or less, then it can be used for the opening CCV of the next 12-hour period.		
	Note: When a required closing CCV of an Aroclor other than Aroclor 1016/1260 is preceded by an opening CCV of Aroclor 1016/1260 CS3 only, the percent difference of each Aroclor peak and surrogate compound must not exceed ± 15%."		
Aro-Item 22	The following Section:		
Exhibit D – Aroclor: Section 9.3.6.7	"If a successful instrument blank and Aroclor 1016/1260 standard cannot be run after an interruption in analysis (Section 9.3.2.6), an acceptable initial calibration must be run before sample data may be collected. All acceptable sample (including LCS and MS/MSDs) and required blank (method/sulfur cleanup) analyses must be preceded and followed by acceptable standards and instrument blanks, as described in Section 9.3.2."		
	is updated to: "If a successful instrument blank and Aroclor 1016/1260 standard cannot be run after an interruption in analysis (Section 9.3.2.6), an acceptable initial calibration must be run before sample data may be collected. All acceptable sample (including LCS and MS/MSDs) and required blank (method/sulfur cleanup) analyses must be preceded and followed by acceptable instrument blanks and standards (opening and closing CCV) as described in Section 9.3.2."		
Aro-Item 23	The following Section:		
Exhibit D - Aroclor: Section 10.2.2.3.1	"Using a syringe or a volumetric pipet, transfer all of the hexane extract to a 10mL vial and, in a fume hood, carefully add 5mL of the 1:1 (v/v) sulfuric acid/water solution." is updated to:		
	"Using a syringe or a volumetric pipet, transfer an aliquot (1 or 2 mL) of the hexane extract to a 10mL vial and, in a fume hood, carefully add 5mL of the 1:1 (v/v) sulfuric acid/water solution."		
Aro-Item 24	The following Sections will be switched:		
Exhibit D – Aroclor: Section 10.2.2.3.1 and 10.2.2.3.2	The language for the updated sentence of Section 10.2.2.3.1 will become Section 10.2.2.3.2 and vice versa.		

HRS Reference #81

EXHIBIT/SECTION(S)	· .	MODIFICATION	N (S)
Aro-Item 25	The following Se	ection:	
Exhibit D – Aroclor: Section 10.3.2.1	"Analytical Sequ	ence	
	All acceptable samples must be analyzed within a valid analysis sequence as given below:		vithin a valid analysis
	Time	Injection #	Material Injected First 12 steps of the initial
	0 hr.	1-12 13	calibration Instrument blank
		14	Aroclor 1016/1260 Standard Sample
	12 hr.	1 st injection past 12 hr.	Last sample Instrument blank
		2 nd injection past 12 hr.	Aroclor 1016/1260 standard Subsequent samples
	Another 12 hrs.		Last sample
		1 st injection past 12 hr.	Instrument blank Aroclor 1016/1260
	:	2 nd injection past 12 hr.	standard
	is updated to:	3 rd injection past 12 hr.	Sample
	"Analytical Sequ All acceptable sa sequence as give	umples must be analyzed v	vithin a valid analysis
	Time	Injection #	Material Injected
	0 hr.	1-12 (or 5-points of all Aroclors) 13 14 15	First 12 steps of theinitial calibration (or 5-points of all Aroclors) Instrument blank Aroclor 1016/1260 Standard Additional Aroclor CS3
	12 hr.	16	Standard (optional) Subsequent Samples Last sample
	·	1 st injection past 12 hr. 2 nd injection past 12 hr.	Instrument blank Aroclor 1016/1260 Standard Detected Aroclor CS3
		3 rd injection past 12 hr.	Standard (as required) Detected Aroclor CS3
	14 hr.	4 th injection past 12 hr.	Standard (as required) Subsequent Samples
	Another 12 hrs.		Last sample
	III.S.	1 st injection past 12 hr.	Instrument blank
		• •	
		3 rd injection past 12 hr.	Sample
		2 nd injection past 12 hr.	Aroclor 1016/1260 standard

EXHIBIT/SECTION(S)

Aro-Item 26

Exhibit D – Aroclor: Section 10.3.2.1.1

MODIFICATION (S)

The following Section:

"The first 12 hours are counted from injection #13, not from injection #1. Samples may be injected until 12:00 hours have elapsed. All subsequent 12-hour periods are timed from the injection of the instrument blank that brackets the front end of the samples. If more than 12 hours elapse between the injections of two instrument blanks that bracket a 12-hour period in which samples or required blanks are analyzed, then the time between the injection of the instrument blank and the preceding sample may not exceed the length of one chromatographic run. While the 12-hour period may not be exceeded, the laboratory may run instrument blanks and standards more frequently, for instance, to accommodate staff working on 8-hour shifts. No more than 14 hours may elapse from the injection beginning the opening CCV (instrument blank) and the injection ending the closing CCV (Aroclor Standard)."

is updated to:

"Injections #1 through #12 in Section 10.3.2.1 may be expanded to include all injections of initial calibration standards as specified in Option 2 and 3 in Section 9.2.3.5. The first 12 hours are counted from injection #13, not from injection #1, in the initial calibration sequence Option 1 detailed in Section 10.3.2.1. Alternately, the first 12 hours will be counted from the injection of the instrument blank of an opening CCV when performed immediately after completion of the initial calibration Options 2 and 3. Samples may be injected until 12:00 hours have elapsed. All subsequent 12-hour periods are timed from the injection of the instrument blank that brackets the front end of the samples. If more than 12 hours elapse between the injections of two instrument blanks that bracket a 12-hour period in which samples or required blanks are analyzed, then the time between the injection of the instrument blank and the preceding sample may not exceed the length of one chromatographic run. While the 12hour period may not be exceeded, the laboratory may run instrument blanks and standards more frequently, for instance, to accommodate staff working on 8-hour shifts. No more than 14 hours may elapse from the injection beginning the opening CCV (instrument blank) and the injection ending the closing CCV (Aroclor Standard)."

EXHIBIT/SECTION(S)	MODIFICATION (S)	
Aro-Item 27	The following:	
Exhibit D – Aroclor: Section 10.3.3.2	"If the response of the largest peak for any Aroclor is greater than the response of the same peak in the high-point standard in the initial calibration for both columns, then the sample must be diluted to have the response of the largest peak of the lower of the two column analyses be between the low and high calibration standards."	
	Is updated to:	
	"If the concentration of the largest peak for any Aroclor is greater than the concentration of the same peak in the high-point standard in the initial calibration for both columns (the largest peak on the second column may be a different peak), then the sample must be diluted to have the concentration of the largest peak of the lower of the two column analyses be between the low and high calibration standards."	
Aro-Item 28	The following:	
Exhibit D – Aroclor: Section 10.3.3.8	"Use the results of the original analysis to determine the approximate DF required to get the largest analyte peak (for the lower of the two column <i>responses</i>) within the initial calibration range."	
	Is updated to:	
	"Use the results of the original analysis to determine the approximate DF required to get the largest analyte peak (for the lower of the two column <i>concentrations</i>) within the initial calibration range."	
Aro-Item 29	The following Section:	
Exhibit D – Aroclor: Section 11.1.1.4	"When an Aroclor other than 1016 or 1260 is detected in a sample, a valid five-point calibration curve specific to that Aroclor must be run, followed by reanalysis of the sample or appropriately diluted sample with the detected Aroclor present. The Mean Calibration Factor (CF) will be used to quantitate the analyte in the sample."	
	is updated to:	
	"When an Aroclor other than 1016 or 1260 is detected in a sample, using a single point calibration, a valid five point calibration of the specific Aroclor must be performed, followed by reanalysis of the sample or appropriately diluted sample (if the sample concentration of Aroclor exceeded calibration) with the Aroclor detected initially. If a valid five-point calibration curve is available for an Aroclor	
	other than 1016 or 1260, the Mean Calibration Factor (\overline{CF}) will be used for quantitation of the Aroclor in the sample, however, quantitation of the surrogate compounds using surrogate data from the initial five-point Aroclor 1016/1260 or from Aroclor 1016 if analyzed as a separate mixture.	
	Note: An estimated concentration (reported with an "S" flag) of the initial detection for an Aroclor other than 1016 or 1260, using a single point calibration standard will be quantitated using the Calibration Factor (CF), of at least 3 major peaks, from the specific single point calibration standard. The surrogates will be quantitated using the initial five-point Aroclor 1016/1260 or from Aroclor 1016 if analyzed as a separate mixture.	

EXHIBIT/SECTION(S)

MODIFICATION (S)

Aro-Item 30

Exhibit D - Aroclor: Section 11.2.1.1.1, Equation 7

The equation is further expanded to allow for greater flexibility in the preparation and cleanup steps as follows:

Concentration
$$\mu g/L = \left(\frac{A_x}{\overline{CF}}\right) \left(\frac{DF}{V_i}\right) \left(\frac{V_t}{V_o}\right) \left(\frac{CV_{out}}{CV_{in} \times E}\right)_1 \left(\frac{CV_{out}}{CV_{in} \times E}\right)_2 \dots \left(\frac{CV_{out}}{CV_{in} \times E}\right)_n$$

where,

A_x = Peak area or peak height of the compound to be measured.

CF = Mean Calibration Factor determined from the initial calibration for the compound to be measured, in area/ng.

DF = Dilution Factor.

V_i = Volume of extract injected in μL.

V_t = Volume of extract produced by the preparation process (extraction and concentration), and before cleanup, in μL.

 V_0 = Volume of the original water sample extracted in mL. Note: for instrument blanks and sulfur blanks assume a volume of $1000 \, mL$.

CV_{out} = Volume of extract produced by a cleanup process (cleanup and concentration), in μL.

 CV_{in} = Volume of extract subjected to a cleanup process, in μL .

E = The efficiency of the cleanup process expressed as a fraction of material that passes through or is not mechanically lost during the cleanup step (e.g. 50% efficiency must be expressed as 0.50)

Aro-Item 31

Exhibit D – Aroclor: Section 11.2.1.2.1, Equation 9

The equation is further expanded to allow for greater flexibility in the preparation and cleanup steps as follows:

Concentration
$$\mu g/kg = \left(\frac{A_x}{\overline{CF}}\right) \left(\frac{DF}{V_i}\right) \left(\frac{V_t}{W_t \times D}\right) \left(\frac{CV_{out}}{CV_{in} \times E}\right) \left(\frac{CV_{out}}{CV_{in} \times E}\right)_1 \cdots \left(\frac{CV_{out}}{CV_{in} \times E}\right)_n$$

where.

 A_{xy} $\overline{\mathrm{CF}}$, DF , $V_{i_1}V_{b_2}CV_{out}$ CV_{in} , and E are the same as Equation 7 above.

 W_t = Weight of the original soil sample extracted in g.

$$D = \frac{100 - \% \text{Moisture}}{100}$$

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 32	The following Section:
Exhibit D – Aroclor: Section 11.2.2	"Target Compounds
	The quantitation of Aroclors must be accomplished by comparing the heights or the areas of each of a minimum of 3 major peaks of the Aroclor in the sample with the $\overline{\text{CF}}$ for the same peaks established during the specific five-point calibration. The concentration of multi-component analytes is calculated by using Equations 7 and 9, where A_x is the area for each of the major peaks of the Aroclor. The concentration of each peak is determined and then a mean concentration for a minimum of 3 major peaks is determined on each column."
	is updated to: "Target Compounds
	Except for an estimated value reported for an Aroclor other than 1016 or 1260, The quantitation of Aroclors must be accomplished by comparing the heights or the areas of each of a minimum of 3 major peaks of the Aroclor in the sample with the CF for the same peaks established during the specific five-point calibration. The concentration of multi-component analytes is calculated by using Equations 7 and 9, where A _x is the area for each of the major peaks of the Aroclor. The concentration of each peak is determined and then a mean concentration for a minimum of 3 major peaks is determined on each column."
Aro-Item 33	The following Section:
Exhibit D – Aroclor: Section 11.2.2.1	"Note that the CFs used for the quantitation of Aroclors are the CFs from the concentration of the specific five-point calibration."
	is updated to: "To quantitate and report the estimated concentration of an Aroclor other than 1016 or 1260, use the Calibration Factor (CF) for a minimum of 3 major peaks, from the single point Aroclor calibration standard used for the Aroclor pattern recognition. It will be necessary to substitute the single Calibration Factor (CF) for the Mean CF (\overline{CF}) in Equations 7, 8, 9 and 10. Note: The CFs used for the quantitation of target Aroclors are the CFs from the concentration of the specific five-point calibration."

EXHIBIT/SECTION(S)

MODIFICATION (S)

Aro-Item 34

Exhibit D - Aroclor: Section 11.2.3.1, Equation 12

The equation is further expanded to allow for greater flexibility in the preparation and cleanup steps as follows:

EQ. 12 Adjusted CRQL Calculation for Water Samples

$$Adjusted CRQL = \left(Contract CRQL\right) \left(\frac{V_x}{V_o}\right) \left(\frac{V_t}{V_y}\right) \left(DF\right) \left(\frac{CV_{out}}{CV_{in} \times E}\right)_1 \left(\frac{CV_{out}}{CV_{in} \times E}\right)_2 \dots \left(\frac{CV_{out}}{CV_{in} \times E}\right)_n$$

where,

Contract CRQL = The CRQL value reported in Exhibit C - Aroclors (µg/L).

 V_x = Contract sample volume (1000 mL).

V₀ = Volume of water extracted in mL. Note: for instrument and sulfur blanks assume a volume of 1000mL.

 V_t = Volume of water concentrated extract in μL .

 V_y = Contract concentrated extract volume (10,000 μ L).

DF = Dilution Factor.

 CV_{out} . = Volume of extract produced by a cleanup process (cleanup and concentration), in μL .

CV_{in} = Volume of extract subjected to a cleanup process, in μL.

E = The efficiency of the cleanup process expressed as a fraction of material that passes through or is not mechanically lost during the cleanup step (e.g. 50% efficiency must be expressed as 0.50).

MODIFICATION (S)

Aro-Item 35

Exhibit D – Aroclor: Section 11.2.3.2 Equation 13

The equation is further expanded to allow for greater flexibility in the preparation and cleanup steps as follows:

EQ. 13 Adjusted CRQL Calculation for Soil/Sediment Samples

$$Adjusted CRQL = \left(Contract CRQL\right) \left(\frac{W_x}{W_s \times D}\right) \left(\frac{V_t}{V_y}\right) \left(DF\right) \left(\frac{CV_{out}}{CV_{in} \times E}\right)_1 \left(\frac{CV_{out}}{CV_{in} \times E}\right)_2 \dots \left(\frac{CV_{out}}{CV_{in} \times E}\right)_n$$

where,

Contract CRQL = The CRQL value reported in Exhibit C - Aroclors (µg/Kg).

 W_x = Contract sample weight (30 g).

 W_s = Weight of sample extracted in grams (g).

 $\mathbf{p} = \frac{100 - \%\text{Moisture}}{100}$

 V_t = Volume of the concentrated extract in μL .

 V_y = Contract concentrated extract volume (10,000 μ L).

DF = Dilution Factor.

 CV_{out} = Volume of extract produced by a cleanup process (cleanup and concentration), in μ L.

 CV_{in} = Volume of extract subjected to a cleanup process, in μL .

E = The efficiency of the cleanup process expressed as a fraction of material that passes through or is not mechanically lost during the cleanup step (e.g. 50% efficiency must be expressed as 0.50).

Aro-Item 36

Exhibit D - Aroclor: Section

11.2.4

The following Section:

"The concentrations for surrogate compounds can be calculated by using Equation 7 (for waters) and Equation 9 (for soils) and the CF from the most recent initial calibration."

is updated to:

"The concentrations for surrogate compounds can be calculated by using Equation 7 (for waters) and Equation 9 (for soils) and the CF from a valid initial five-point calibration of Aroclor 1016/1260 or from Aroclor 1016 if analyzed as a separate mixture."

EXHIBIT/SECTION(S)	MODIFICATION (S)
Aro-Item 37	The following Section:
Exhibit D – Aroclor: Section 11.3.5	"The RT for each of the surrogates must be within the RT window (Section 9.2.4.3) for both GC columns."
	is updated to:
	"Surrogate compounds Retention Time (RT) must be compared to the window established during a valid initial five-point calibration of Aroclor 1016/1260 or from Aroclor 1016 if analyzed as a separate mixture. The RT for each of the surrogates must be within the RT window (Section 9.2.4.3) for both GC columns."
Aro-Item 38 Exhibit D – Aroclor: Section 12.3.4.2	The following Section: "Calculate individual compound recoveries of the LCS using Equation 14"
	is updated to: "Calculate individual compound recoveries of the LCS using Equation 15 ".